SOME PHYSIOLOGICAL REACTIONS OF NEW ZEALAND RABBIT TO DRINKING WATER POLLUTED BY THE INSECTICIDE "PROFENOFOS"

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SUMMARY

The present investigation was carried out on New Zealand rabbits to determine the physiological reactions through liver, kidney, and thyroid functions and relevant blood compounds with the polluted water by 75 and 150 ppm of the insecticide Profenofos. The animals were exposed to the polluted water drinking for 60 days. Blood samples were collected at 30 and 60 days. Male rabbit given sub-lethal doses of 75ppm and 150ppm profenofos in drinking water showed a significant increase in the red blood cells count (RBCs), haemoglobin concentration (Hb) and packed cell volume (PCV) at 150ppm of profenofos after 60 days (116.6% & 110.6%, 109.8% & 114% and 116.7% & 112% respectively), a significant elevation in the mean corpuscular volume (MCV) was noted at the highest concentration (150ppm) after 30 days, whereas a significant decrease occurred in the values of mean corpuscular haemoglobin concentration (MCHC) after 60 days, (75.3% and 87.7% respectively). A significant decrease in the total white blood cells (leukopenia) was observed in treated rabbit. There was an elevation in (ALT) activity after 60 days and the increased values was 124.3% for 150ppm relative to control, also a decreases in concentration of albumin (ALB) was observed. The Profenofos caused inhibition of cholinesterase activity almost at all the treatment periods. There was not any marked change in cholesterol concentration throughout the experimental period (60 days). Also the level of the urea in treated rabbits decreased after 30 days of treatment with tested concentration 43.7% & 34.2% for 75 and 150 ppm respectively. The creatinine concentration decreased significantly after 60 day from treatment with high treatment (150 ppm).

Keywords: Profenofos insecticide, polluted water, liver, kidney, thyroid functions, hematology

INTRODUCTION

As the demand for agricultural products increases, so inevitably does the need for pesticides. Currently organophosphorus compounds are part of a large group of
synthetic pesticides that have been developed in the last 50 years, and are commonly used in agricultural and public health. Due to their wide use, contamination of food, water, and air has become imminent and consequently adverse health effects are inevitable in humans, animals, wildlife and fish (Gupta, 1994). The use of pesticides not only increased the crop yields but also helped in controlling ectoparasites on livestock. However, widespread use of these chemicals is not without risks. There have been several instances of poisoning in animals. Animals get exposed to these pesticides due to accidental ingestion due to their inadvertent use for the control of ectoparasites and insects in the vicinity of poultry, rabbit, and animal houses. Residues of the grain protectant pesticides may also come in animal feeds. Perusal of literature did not reveal the toxicity of "Profenofos" insecticide on rabbit in Egypt.

The aim of this study is to determine the effect of polluted water by profenofos on liver; kidney and thyroid gland functions and on the changes in relevant blood compounds as physiological reactions in male New Zealand rabbits as farm animals.

MATERIALS AND METHODS

Experimental animals:

New Zealand rabbits weighing 2.4±0.3 kg were supplied by the farm of General Organization of Serum and Vaccine (Helwan farm). The present studies were performed on 30 mature male New Zealand rabbits. The animals were allowed to acclimatize to laboratory conditions for a period of 3 weeks prior to the experiment. The animals were kept on balanced diet throughout the experimental period. Feed and water were available ad lib during the experiment.

Experimental design:

Randomized groups, of these rabbits housed in cages, were allocated into three treatment groups, (10 each) and treated with Profenofos through drinking water for 60 successive days. The animals groups were as follow: group (A) rabbits kept without any treatment (as a control), and Groups (B) and (C) rabbits treated with Profenofos in drinking water at 75 and 150 ppm, respectively. On day 30 and 60, blood samples were taken from each rabbit.

Blood samples:

Blood samples were collected from Occuseal veins by heparinized capillary tubes at into clean, dry and labeled eppendorf of tubes (1.5 ml). Each blood sample was divided into two aliquots, one containing EDTA (Ethylendiamine tetra acet acid) as anticoagulant (1mg/ml blood) for studying the hemogram of the experimental animals, the second aliquot was left at room temperature for 2 hours before centrifugation at 3500 rpm. for 15 minutes in a refrigerated centrifuge to separate serum, and kept in a deep freezer at (-20 °C). Serum samples were examined for studied parameters. Counts of red and white blood cells, hemoglobin concentration, and packed cell volume were determined according to the methods of Schalm (1986). The erythrocytes indices; mean corpuscular volume (MCV), mean corpuscular haemoglobin (MCH) and mean corpuscular haemoglobin concentration (MCHC) were also calculated depending on RBCs count, haemoglobin concentrations (Hb) and packed cell volume (PCV) as follow:

\[
MCV = \left\{ \frac{PCV \ (vol \%) \times RBCs \ (million/ml)}{PCV} \right\} \times 10 = F1
\]
Thyroxin (T₄), and Tri-iodothyronine (T₃) were determined in serum according to Britton et al. (1975) by the coat-A-count technique, using kits purchased from Diagnostic Products Corporation (DPC).

The following physiological parameters expressed the liver function:

- The activities of alanine aminotransferase (ALT) and aspartate aminotransferase (AST) were determined spectrophotometrically according to Reitman and Frankel (1957).
- Serum total protein concentration (T.P g/dl) was determined spectrophotometrically according to Henry et al. (1974).
- Serum albumin concentration (ALB g/dl) was determined spectrophotometrically by the method of Bartholomew and Delaney (1964).
- Alkaline phosphatase activity (ALP U/l) was colorimetrically estimated according to the method of Empfehlungen (1972).
- The activity of cholinesterase (ChE) was determined kinetically according to the method of Waber et al. (1966).
- Serum cholesterol concentration was determined spectrophotometrically by the method adopted by Siest et al. (1981).

The following physiological parameters are expressed the kidney function: Serum urea concentration (g/dl) was measured spectrophotometrically by the method of Patton and Crouch (1977). Serum creatinine (mg/dl) was measured spectrophotometrically by the method of Faulkner and King (1976).

**Statistical analysis**

Data were subjected to statistical analysis by least squares analysis of variance 2×2 factorial design. The main effects of treatment and duration and all possible interaction terms were included in the model according to the procedures reported by Snedecor and Cochran (1980); and "t" test was performed to evaluate the difference between mean values of the treated groups and those of the control group.

**RESULTS AND DISCUSSION**

Effects of Profenofos on haematological parameters and T₃ and T₄ of rabbits are shown in Table (1). Male rabbits given sub-lethal doses of 75ppm and 150ppm Profenofos in drinking water showed a significant increase in the red blood cells count (RBCs), haemoglobin concentration (Hb) and packed cell volume (PCV), being significant at 150ppm. The percentage increases by 150 ppm after 30 and 60 days were (16.6% & 17.6% & 5.8% and 15%; 17.7% & 13% respectively). The exposures of rabbit to both doses of Profenofos lead to a significant decrease in the white blood cells (WBCs) after 30 days the decrease with percentages of 21.3% for 75 ppm and 11.5% for 150 ppm relative to control. But WBCs increased significantly after 60 days for 150 ppm. In the present study, a significant elevation in the mean corpuscular volume (MCV) was noted by the highest treatment (150ppm) after 30 days. It is clear that the level of MCV after 60 days was lower than after 30 days, and the differences between MCV values at 75 ppm and 150 ppm were not significant. However a significant decrease in the values of mean corpuscular haemoglobin (MCH) and mean corpuscular haemoglobin concentration (MCHC) were recorded after 60 days, (75.3% and 87.7% respectively).
Table 1. Effect of Profenofos treatments on haemogram parameters in male New Zealand rabbit

<table>
<thead>
<tr>
<th>Dose Parameters</th>
<th>Control</th>
<th>75 ppm</th>
<th>150 ppm</th>
<th>Control</th>
<th>75 ppm</th>
<th>150 ppm</th>
</tr>
</thead>
<tbody>
<tr>
<td>RBCs $\times 10^6$ μl</td>
<td>6.90±0.1</td>
<td>7.00±0.27</td>
<td>8.11±0.23*</td>
<td>7.18±0.12</td>
<td>7.86±0.25</td>
<td>8.43±0.09**</td>
</tr>
<tr>
<td>WBCs $\times 10^3$ μl</td>
<td>6.1±0.71</td>
<td>4.5±0.36*</td>
<td>5.40±0.33*</td>
<td>7.4±0.73</td>
<td>6.76±0.17</td>
<td>9.12±0.98*</td>
</tr>
<tr>
<td>Hb g/dl</td>
<td>16.11±1.24</td>
<td>18.81±0.66</td>
<td>17.02±1.15*</td>
<td>16.90±0.34</td>
<td>15.01±0.32</td>
<td>19.43±0.16**</td>
</tr>
<tr>
<td>PCV (%)</td>
<td>51.5±2.63</td>
<td>52.5±1.32</td>
<td>64.3±1.48**</td>
<td>51.00±0.83</td>
<td>55.00±0.63</td>
<td>58.7±1.42**</td>
</tr>
<tr>
<td>MCV Fl</td>
<td>70.4±2.8</td>
<td>75.96±2.4</td>
<td>90.6±0.9**</td>
<td>66.49±0.23</td>
<td>68.05±3.1</td>
<td>70.46±2.5</td>
</tr>
<tr>
<td>MCH Pg</td>
<td>21.20±1.1</td>
<td>19.81±0.33</td>
<td>22.67±1.19</td>
<td>20.37±1.79</td>
<td>23.53±1.01</td>
<td>18.88±0.34*</td>
</tr>
<tr>
<td>MCHC (%)</td>
<td>33.4±3.33</td>
<td>28.02±2.0</td>
<td>29.31±1.94</td>
<td>32.9±0.7</td>
<td>29.04±0.6**</td>
<td>25.7±2.05*</td>
</tr>
<tr>
<td>T₄ mol/l</td>
<td>68.00±2.7</td>
<td>62.55±1.2</td>
<td>65.38±5.55</td>
<td>75.52±3.74</td>
<td>60.32±6.31</td>
<td>54.49±3.35**</td>
</tr>
<tr>
<td>T₃ mol/l</td>
<td>1.133±0.1</td>
<td>1.32±0.03</td>
<td>1.52±0.1</td>
<td>0.825±0.12</td>
<td>1.24±0.07*</td>
<td>11.95±0.17**</td>
</tr>
</tbody>
</table>

Each value represents mean ± SE. **Significant differences versus control at p < 0.01
*Significant differences versus control at p < 0.05

Sub-chronic toxicity by these Profenofos treatment induced erythrocytosis in male rabbits after dosing with 150ppm for 60 days as well as an elevation in the PCV values, this condition referred to absolute polycythemia, which may be caused by hypoxia in treated animals meanwhile, a significant increase in the PCV values in the 150ppm group after 60 days may be attributed to dehydration in the Profenofos, treated rabbits. This data are in accordance with those reported by Jain (1993).

A significant decrease in the total white blood cells (leukopenia) observed in treated rabbit may be attributed to lymphopenia resulting from release of stress hormone (corticosteroids) in treated rabbits as suggested by Marx (1996). The results in Table (1) indicate that male rabbits treated with profenofos showed a significant decrease in the concentration of thyroxin (T₄) and Tri-iodothyronine (T₃) (hypothyroidism) after 30 days of treatment by 150 ppm and after 60 days by both 75 ppm and 150 ppm. The results of this present work are in agreement with those found by Abd Ellah (1987), Farid (1997), Hotz et al. (1997), and Salem et al. (1999). They stated that, a significant decrease of T₄ and T₃ occurred after administration of different pesticides to experimental animals.

Data in Table (2) illustrates that treatment of male New Zealand rabbits with Profenofos resulted in elevation in (ALT) activity after 60 days and the increased values was 24.3% for 150 ppm relative to control, although the enzyme activity was decreased significantly (74%) after 30 days of treatment for the concentration 75 ppm., El-Said (1997) and El-Halwagy (2000, mentioned that ALT and AST did not alter significantly after treatment with Diniconazole and Monocrotophos at all tested dose for 90 days in rats. Hashem (1980) reported that the depression in the activity of transaminases might be due to the formation of complex compounds with ALT or AST in the liver. Aspartate aminotransferase (AST) and alanine aminotransferase (ALT) have long been used as sensitive indicators of liver disease in animals and have been regarded as being virtually liver specific (Wills 1985).

The data in table (2) shows that there is not change in total protein (TP) concentration throughout the experimental period. It should be mentioned that the present data are in a quite good agreement with those reported by El-Kashoury (1999) and Hassan et al. (2002) who reported that total protein concentration changed insignificantly in experimental animals exposed to Dimethoate (OP), Oxamyl (carbamate insecticide), Malathion, Fenitrothion and mixture of some synthetic pesticides (Imidacloprid, Profenofos, and Carbosulfan). On the other hand; Zidan et
al. (1998) and Choudhary & Joshi (2002) reported increase in total protein concentration by treatment with some synthetic pesticide.

Table 2. Effect of Profenofos on liver and kidney functions of male New Zealand Rabbit

<table>
<thead>
<tr>
<th>Dose Parameters</th>
<th>Control</th>
<th>after 30 (days) 75 ppm</th>
<th>150 ppm</th>
<th>Control</th>
<th>after 60 (days) 75 ppm</th>
<th>150 ppm</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALT U/ml</td>
<td>176±11.0</td>
<td>170±13.0*</td>
<td>130.2±23.2</td>
<td>180.6±23.9</td>
<td>190±12.5</td>
<td>223.7±6.0*</td>
</tr>
<tr>
<td>AST U/ml</td>
<td>153±21.5</td>
<td>136.2±4.2</td>
<td>127±24.2</td>
<td>184±23.9</td>
<td>194±7±2.7</td>
<td>173.3±6.6</td>
</tr>
<tr>
<td>T.P g/dl</td>
<td>6.44±0.7</td>
<td>8.13±0.9</td>
<td>7.67±0.8</td>
<td>6.26±0.5</td>
<td>6.59±0.6</td>
<td>6.55±6.8</td>
</tr>
<tr>
<td>ALB g/dl</td>
<td>3.68±0.1</td>
<td>4.02±0.25</td>
<td>3.65±0.1</td>
<td>4.51±0.1</td>
<td>3.72±0.4</td>
<td>3.46±0.1**</td>
</tr>
<tr>
<td>ALP U/l</td>
<td>60.42±3.41</td>
<td>51.79±5.3</td>
<td>37.06±4.98**</td>
<td>68.02±13.14</td>
<td>65.6±7.8</td>
<td>34.1±6.36**</td>
</tr>
<tr>
<td>ChE 600.6±41.6</td>
<td>480.3±67.4</td>
<td>350.3±43.64*</td>
<td>580.7±35.19</td>
<td>397.3±17.4*</td>
<td>444.4±48.9*</td>
<td>397.3±17.4*</td>
</tr>
<tr>
<td>Cholesterol mg/dl</td>
<td>0.713±0.44</td>
<td>0.714±0.10</td>
<td>0.64±0.08</td>
<td>0.647±0.06</td>
<td>0.665±0.1</td>
<td>0.57±0.185</td>
</tr>
<tr>
<td>Urea g/dl</td>
<td>0.173±0.06</td>
<td>0.097±0.04</td>
<td>0.114±0.05**</td>
<td>0.181±0.05</td>
<td>0.079±0.1</td>
<td>0.099±0.06**</td>
</tr>
<tr>
<td>Creatinine mg/dl</td>
<td>1.675±0.22</td>
<td>0.324±0.12**</td>
<td>0.268±0.04**</td>
<td>1.701±0.06</td>
<td>0.651±0.09**</td>
<td>0.548±0.04**</td>
</tr>
</tbody>
</table>

Each value represents mean ± SE

**Significant differences versus control at p < 0.01
*Significant differences versus control at p < 0.05

Data in Table (2) illustrates that; the Profenofos induced significant decrease in concentration of albumin (ALB) after 60 days of treatment with high dose of (76.8% for 150 ppm). The present results are generally similar to those reported by several workers. Ayyat (2000) and Ahmed et al. (2002), who mentioned that total albumin, was altered significantly according to the type of different pesticides.

Data in Table (2) reveal that Profenofos induced significant decreases in activity of alkaline Phosphates (ALP) after 30 and 60 days of treatment under high concentration of profenofos where values of ALP activities reached 45.5% and 51.9%, respectively compared with control. These results were supported by the degenerative changes and necrotic change observed in the liver by the histopathological examination. The present results are in agreement with those reported by El-Kashoury (1999). Who reported that, rats exposed to formulated mixture of three organophosphorus pesticides showed a significant decrease in ALP activities. Hassan et al. (2002) reported that plasma alkaline phosphatase (ALP) did not alter throughout the experimental period (60 days) at 500 mg/kg Bwt of Fenitrothion (OP) in treated New Zealand male rabbits.

The present results revealed that Profenofos caused inhibition of cholinesterase activity almost at the two periods. The results are in agreement with those obtained by Ahmed et al. (2002), Zayed et al. (2003), Rahman & Siddiqui (2003) and Maryam Akhgari et al. (2003), they mentioned that, ChE activity was decreased significantly in the experimental animals after exposure to organophosphorus compounds.

The results in Table (2) revealed that Profenofos administrated to male New Zealand Rabbits at the two tested concentrations did not cause any marked changes in cholesterol concentration throughout the experimental period (60 days).

In mammals, the kidney functions as a major excretory organ for elimination of metabolic wastes from the body. In most species, death occurs within a week after total cessation of renal function. Urea is biosynthesized in the liver from ammonia. Data presented in table (2) revealed a decrease in the level of the urea in treated rabbits after 30 days of treatment by 43.7% & 34.2% for 75 and 150 ppm, respectively. The
results are in agreement with those recorded by El Kashoury (1999) who observed that urea concentration was decreased significantly in Imidaclopride, Profenofos and Carbosulfan mixture treated rats.

Creatinine is a waste product derived from creatine and creatine phosphate. It is clear from data (Table 2) that creatinine decreased significantly after 60 day by treatment with 150 ppm. Values of creatinine concentration reached 39% relative to control. The present results are in accordance with those reported by El-Said (1997). They found that creatinine concentration was decreased in experimental animals administered with synthetic pesticides.

CONCLUSION

The studied criteria included: relevant blood traits as well as, liver, kidney and thyroid functions. The male New Zealand rabbit were treated with Profenofos through drinking water for 60 days with two concentrations (75 and 150 ppm). Profenofos reduced ALT, Cholinesterase (ChE) and alkaline phosphates activities after 30 days at low and high concentrations (75 ppm and 150-ppm). no significant changes appeared after 30 and 60 days in AST activity and in cholesterol and TP concentrations. Albumin (ALB) concentration decreased in rabbits treated by 150 ppm after 60 days. But, the highest concentration (150 ppm) showed significant increase in albumin content after 60 days of treatment. Urea and Creatinine concentrations were decreased significantly by 75 and 150 ppm for 30 days. Profenofos induced a significant increase in the level of T3 and T4 after 60 days at 75 ppm. RBCs counts, haemoglobin concentration and PCV were increased significantly with the different concentrations of Profenofos, but WBCs counts and erythrocyte indices (MCV, MCH and MCHC) were decreased significantly in rabbits with profenofos at 75 and 150 ppm.

REFERENCES


(Cited by Zidan et al. 1998).
الاستجابات الفسيولوجية في الأراب النيوزلندي نتيجة شرب ماء ملوث بمادة البروفينوفوس

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أجري البحث في كلية الزراعة جامعة القاهرة بالجيزة على ثلاثين ذكر أراب من نوع النيوزلندي للструктур على أهم التغيرات الحادة في تركيب الدم وكذا التغير في وظائف الكبد والكلى والرئة كرد فعل فسيولوجي للشرب لمدة 60 يوم ماء ملوث بكميات أقل من الحد المسمى (75 جزء من المليون و 150 جزء من المليون) من مركب البروفينوفوس المستخدم لمكافحة الطفيليات الخارجية للحيوانات المزرعة المختلفة كما أنه يستخدم أيضا لمكافحة الأتاف الحشرية لمنع المحاصيل الزراعية والتي يستخدم معظمها كعلف حيوي.

أولا: التغيرات الحادة في تركيب الدم:

حدث زيادة معنوية لعدد كريات الدم الحمراء بالمعالجة بالجرعة العالية بعد 60 يوم وحدث زيادة معنوية لتركيز الهيموجلوبين في التركيز العالي بعد 60 يوم من المعالمة بالسم بالمركب وكذا حدوث نقص معنوي ملحوظ لعدد كريات الدم البيضاء للتركيز العالي والمنخفض بعد 30 يوم من المعالمة بالسم.

حدث زيادة معنوية لقيمة المكونات الخلوية بالدم (PCV) للتركيز العالي بعد 60 يوم من المعالمة كما حدث زيادة معنوية لقيمة PCV في التركيز المنخفض بعد 60 يوم من المعالمة. وحدث زيادة معنوية لقيمة MCV في التركيز العالي بعد 30 يوم من المعالمة وذلك بالنسبة للتركيز العالي بينما حدث انخفاض معنوي في قيمة MCV في التركيز المنخفض بعد 60 يوم من المعالمة. وحدث انخفاض معنوي للتركيز المنخفض لقيمة MCHC في التركيز المنخفض بعد 60 يوم من المعالمة.

ثانيا: القياسات الفسيولوجية التي تستدعي كدليل على حدوث التغيرات في وظائف الكبد ويمكن إيجازها فيما يلي:

1 حدوث نقص معنوي في نشاط إنزيم ALT بعد 30 يوماً من المعالمة ويدل على حدوث زيادة معنوية ملحوظة.
2 عدم حدوث أي زيادة أو نقص معنوي في نشاط إنزيم AST (الإنزيم تقل مجموعه الأمين).
3 عدم حدوث أي تغيرات معنوية في مستوي البروتين الكلي.
4 حدوث نقص معنوي في مستوي الألبومين خلال مراحل التجربة.
5 انخفاض نشاط مستوي إنزيم الاستييل كولين استيروج خلال مراحل التجربة.
6 حدوث انخفاض في مستوي إنزيم الفسفاتاز الفاقد لخلايا المرحلة الأولى والثانية من المعالمة فقط.
7 عدم حدوث أي اختلافات معنوية في مستوي الكولستيروال خلال مراحل التجربة المختلفة.

ثالثا: حيث حدث نقص معنوي في مستوي البروتينات والكربوهيدرات كدليل على حدوث التغيرات في وظائف الكبد.
رابعًا: التغيّرات في وظائف الهرمونات حيّث:

1 - حدوث انخفاض في تركيز هرمون التريوتروسين (T4) بعد 60 يوم من المعالجة.
2 - حدوث زيادة معنوية لهرمون التريالي أوتيديدين (T3) بعد 60 يوم من المعالجة.