Study of Effect of Methomyl on Some Hematological, Biochemical Parameters and Histological Changes in Male Albino Rats

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Abstract

This study has been designed to study the effect of methomyl on hemodynamic, liver and kidney in rats. Methomyl compound (0.5, 1, and 2 mg/kg. bw) was orally administered once per a day during a period of 28 days of experiment. Results showed a significant (P<0.05) decrease in the level of WBC, HCT, MCHC and a non-significant decrease in the level of RBC, HB, MCV, MCH, was observed in the methomyl group, while the level of PLT significantly increased in Methomyl group in comparison with control group. Non-significant decrease in a level of serum urea in all three groups of Methomyl, while serum creatinine level deceased non-significantly in group1 and group 2, but increased non-significantly in group3 of Methomyl when compared with control group. A Significant decrease in the level of HDL, with significant increase in the level of serum cholesterol, ALT and AST. A non-significant increase in the level of TG, LDL, VLDL and ALP of methomyl group when compared with control group. The Histology of liver of rat treated with 0.5,1 and 2, mg / kg bw. methomyl showed that dilation of central vein, sinusoids between hypertrophied hepatocytes and cytoplasmic vacuolization with loss of radial arrangement of cells, Meanwhile, kidney showed some changes including glomerular degeneration, tubular degeneration, hemorrhage, infiltration and glomerular shrinkage.

Keywords: Methomyl, hematological,biochemical change, lipid profile, histopathology.
تأثير الميثوميل على بعض الصفات الدموية والكيميائية والتغير النسجي في ذكور الجرذان البيض

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الخلاصة

صممت هذه التجربة لدراسة تأثير مبيد الميثوميل على بعض مكونات الدم والكبد والكلية في الجرذ. الجرعات المستخدمة كانت (0.5 و 1 و 2 ملغ/كغم وزن الجسم). تم تجريع الجرذان عن طريق الفم يوميا لمدة 28 يوما. أشارت النتائج أن تأثير الميثوميل على بعض مكونات الدم والإنتاج الكيماوي والكبد والكلي في الجرذ مثل ما ورد في الأدب العلمي. وجدت الإحصاءات أن هناك تأثيراً غير معنوي في مستوى كريات الدم البيض وحجم خلايا الدم المتراصة ومعتدل تكاثر خضاب الدم والانخفاض غير معنوي في مستوى كريات الدم الحمر بالنسبة خضاب الدم ومعتدل حجم الخلايا ومعتدل خضاب الدم في مجموعة الميثوميل. ورتفعت مستوى الصفائح الدموية في مجموعة الميثوميل عند مقارنتها بمجموعة التحكم. بينما النتائج أن تأثير الميثوميل على بعض مكونات الدم والإنتاج الكيماوي والكبد والكلي في الجرذ مثل ما ورد في الأدب العلمي. وجدت الإحصاءات أن هناك تأثيراً غير معنوي في مستوى كريات الدم البيض وحجم خلايا الدم المتراصة ومعتدل تكاثر خضاب الدم والانخفاض غير معنوي في مستوى كريات الدم الحمر بالنسبة خضاب الدم ومعتدل حجم الخلايا ومعتدل خضاب الدم في مجموعة الميثوميل. ورتفعت مستوى الصفائح الدموية في مجموعة الميثوميل عند مقارنتها بمجموعة التحكم. ووجدت الإحصاءات أن هناك تأثيراً غير معنوي في مستوى كريات الدم البيض وحجم خلايا الدم المتراصة ومعتدل تكاثر خضاب الدم والانخفاض غير معنوي في مستوى كريات الدم الحمر بالنسبة خضاب الدم ومعتدل حجم الخلايا ومعتدل خضاب الدم في مجموعة الميثوميل. ورتفعت مستوى الصفائح الدموية في مجموعة الميثوميل عند مقارنتها بمجموعة التحكم. ووجدت الإحصاءات أن هناك تأثيراً غير معنوي في مستوى كريات الدم البيض وحجم خلايا الدم المتراصة ومعتدل تكاثر خضاب الدم والانخفاض غير معنوي في مستوى كريات الدم الحمر بالنسبة خضاب الدم ومعتدل حجم الخلايا ومعتدل خضاب الدم في مجموعة الميثوميل. ورتفعت مستوى الصفائح الدموية في مجموعة الميثوميل عند مقارنتها بمجموعة التحكم.

الكلمات المفتاحية:
الميثوميل، التغير الدموي والكيميائي، صورة الدهون، التغير النسجي

Introduction

People in the life exposed to an environmental stressors as a routine part of their existence, about 95% of all chemical toxicity studies were implement on individual chemicals (1, 2). The biological activity of a chemical may be readjustment through concurrent exposure of a test organism to another chemical agent and such interactions might result in addition, or reduction of the effect of the chemical (3). Methomyl S-methyl N-(methylcarbamoyloxy) thioacetamide
Methomyl (C₃H₁₀N₂O₂S), is one of the most common pesticides which are used in the control of insects. It is used in health programs and agriculture; it causes many toxic effects (4). Animals and human exposure occur during spraying of insecticide and ingestion of food contaminated with methomyl (5). The exposure to this insecticide exerts neurodegenerative with high mortality rates (6). Methomyl insecticides inhibit acetyl cholinesterase enzyme activity in the central and peripheral nervous system when ingested in acute exposure (7). Many authors showed that these compounds in both acute and chronic intoxication disturb the redox processes, changing the activities of antioxidative enzymes and causing enhancement of lipid peroxidation in many organs in the body (8). Methomyl induced toxicity against the treated rats (9), by exerting its toxic effect via peroxidative damage to the liver, kidney and splenic cell membranes and cause DNA damage in these organs (10). The changes in biochemical parameters as measured in various body fluids may be the more sensitive indicators due to exposure to pesticides in the environment (11), and histopathological changes consider as a marker for environmental stress when these pesticides used for a long time (12). The aim of the present study is to assess the effect of methomyl on some hematological, kidney and liver function test with lipid profile parameters and histological changes in treated rat’s organ.

Materials and Methods

Chemicals
Methomyl (Methomex, S-methyl N-(methylcarbamoyloxy) thioacetamide was obtained from the Central Laboratories of Agricultural Pesticides, as a pure white very small crystal powder, it was dissolved in saline and prepared freshly.

Animals And Housing
Twenty four adult male albino rats were used, the weight of rats were about 200 -250 g . Animals were housed in plastic cages bedded with wooden chips. They were housed under standard laboratory conditions, 12:12 light/dark photoperiod at 22±2 ºC. The animals were given tap water and standard rat pellets ad libitum.
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Experimental Design
The rats were divided into four groups, each with six animals and continued for 28 days as follows:
Group 1: Control. The rats were given standard rat chow and ad libitum and 0.5 ml of saline orally.
Group 2: The rats were given standard rat chow with Methomyl by oral gavage (0.5 mg/kg bw) daily
Group 3: The rats were supplied with standard rat chow with Methomyl by oral gavage (1 mg/kg bw) daily
Group 4: The rats were supplied with standard rat chow with Methomyl by oral gavage (2 mg/kg bw) daily

Body and Organs Weight
Initial and final body weights were recorded for the calculation of body weight gain, rats were dissected and the liver and kidney weight were determined by using sensitive balance.

Blood Sampling
Ketamine hydrochloride (100mg/ kg bw) used and 5ml of blood samples drawn from the heart puncture for each rat in each group, 2ml of blood collected into heparinized tubes for determination of some hematological parameters such as white blood cell (WBC) count, red blood cell (RBC) count, hemoglobin (Hb) concentration, hematocrite (HCT) level, red blood cell indices Mean corpuscular hemoglobin(MCH), Mean corpuscular volume( MCV) and Mean corpuscular hemoglobin concentration(MCHC) and platelet (PLT) count by using coulter counter (Japan). Other 3ml of blood collected in nonheparinized tubes then centrifuged at 3000 rpm for 15 minutes, the serum was used for determination of biochemical parameters like serum urea, creatinine, lipid profile such as cholesterol, Triglycerides( TG), High density lipoproteins( HDL), Low density lipoprotein(LDL) and Very low density lipoprotein(VLDL), liver function
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test Alanine transaminase (ALT), Aspartate transaminase (AST) and Alkaline phosphatase (ALP) using kit for each test by Cobas E411.

Histological Examination

The kidney and liver were removed and fixed in 10% neutral buffered formalin solution, after preparation the sections were stained with hematoxylin and eosin (13) and examined using a light microscope.

Statistical Analysis

Data were expressed as mean ± standard error (SE), and were analyzed using Statistical Package for Social Science (SPSS/ Version 20) software. Significance between experimental groups was determined using one way analysis of variance (ANOVA) followed by least significant difference (LSD) test for comparison between two groups. P values less than 0.05 were considered statistically significant.

Results

There was no death during the experimental course (28 days) of the present study. Table (1) shows significant decrease in WBC count (9.233±0.234) in G1, (9.200±0.829) in G2 and (9.583±0.224) in G3 when compared with control group (12.400±0.436), RBC count in G1 was (6.878±0.126), G2 (6.772±0.204) and G3 (6.917±0.183) and Hb level in G1 was (13.262±0.122), G2 (13.150±0.226) and G3 (13.067±0.271) which was decreased non significantly when compared with control group for RBC count was (7.163±0.116) and Hb level was (13.717±0.079) respectively. Level of HCT was decreased significantly in G1 (36.733±0.684) and G3 (36.800±0.301) and increased significantly in G2 (37.900±0.323) compared with control group (37.750±0.224). MCV was decreased non-significantly in methomyl group which were (53.17±0.477) in G1, (53.17±0.601) in G2 and (51.50±0.342) in G3 when compared with control group (53.50±0.563). MCH was decreased non-significantly in G1 (18.50±0.224) and G3 (18.67±0.211) and increased non-significantly in G2 (19.17±0.167) when compared with control group (19.00±0.365). While MCHC was decreased significantly
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in G1, G2 and G3 (35.33±0.715), (35.83±0.307) and (35.67±0.422) respectively when compared with control group (38.17±0.307). Significant increase observed in PLT count in G1 (275.83±42.040) and G3 (229.50±25.947), and decreased in G2 (155.17±8.392) methomyl in comparison with control group (161.83±7.011).

Table (1): Effect of Methomyl on Some Hematological Parameters in Male Rats and Control Group after 28 days.

<table>
<thead>
<tr>
<th>Group</th>
<th>Control (0.5 mg/kg bw)</th>
<th>G1 (1 mg/kg bw)</th>
<th>G2 (2 mg/kg bw)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>WBC×10⁶/mm³</td>
<td>12.400 ±0.436</td>
<td>9.233 ±0.234</td>
<td>9.200 ±0.829</td>
<td>9.583 ±0.224</td>
</tr>
<tr>
<td>RBC×10⁶/mm³</td>
<td>7.163 ±0.116</td>
<td>6.878 ±0.126</td>
<td>6.772 ±0.204</td>
<td>6.917 ±0.183</td>
</tr>
<tr>
<td>Hb g/dl</td>
<td>13.717 ±0.079</td>
<td>13.262 ±0.122</td>
<td>13.150 ±0.226</td>
<td>13.067 ±0.271</td>
</tr>
<tr>
<td>HCT%</td>
<td>37.750 ±0.224</td>
<td>36.733 ±0.684</td>
<td>37.900 ±0.323</td>
<td>36.800 ±0.301</td>
</tr>
<tr>
<td>MCV Fl</td>
<td>53.50 ±0.563</td>
<td>53.17 ±0.477</td>
<td>53.17 ±0.601</td>
<td>51.50 ±0.342</td>
</tr>
<tr>
<td>MCH pg</td>
<td>19.00 ±0.365</td>
<td>18.50 ±0.224</td>
<td>19.17 ±0.167</td>
<td>18.67 ±0.211</td>
</tr>
<tr>
<td>MCHC %</td>
<td>38.17 ±0.307</td>
<td>35.33 ±0.715</td>
<td>35.83 ±0.307</td>
<td>35.67 ±0.422</td>
</tr>
<tr>
<td>PLT×10⁵/mm³</td>
<td>161.83 ±7.011</td>
<td>275.83 ±42.040</td>
<td>155.17 ±8.392</td>
<td>229.50 ±25.947</td>
</tr>
</tbody>
</table>

Table(2) exhibits the serum concentration of urea, creatinine, Cholesterol, TG, HDL, LDL, VLDL, ALT, AST and ALP after 28 days of the oral administration of Methomyl at three different dose levels. According to the kidney function test, serum urea was decreased non-significantly in all three groups which was (20.350±0.888) in G1, (18.416±0.808) in G2 and (17.600±0.524) in G3 in comparison with control group (21.261±0.299). Serum creatinine was decreased non-significantly in G1 and G2 which were (0.480±0.022) and (0.483±0.01) in G1 and G2 respectively, but increased non-significantly in G3(0.545±0.015) when compared with control group (0.540±0.025). Significant increase were observed in the concentration of Cholesterol in all three groups of Methomyl (65.500±1.335, 79.000±1.861 and 69.833±3.590) in G1, G2 and G3 respectively in comparison with control group(62.000±0.862), and non-significant increase in triglyceride concentration which was (41.166±4.206) in G1,
(49.833±5.940) in G2 and (43.000±4.618) in G3 when compared to control group value (34.333±3.382). HDL decreased significantly, in G1 was (24.133±0.717), in G2 was(24.567±1.322) and in G3 was (20.933±0.581) in comparison with control group which was (28.900±2.487). LDL, VLDL these two parameters increased non-significantly in Methomyl groups, about LDL in G1 was(31.666±1.666), in G2 was (44.266±1.150), in G3 was (38.666±2.691), VLDL in G1 was (8.333±0.881), in G2 was(8.500±1.118) and in G3 was(7.833±0.872) when compared with control group of LDL (26.833±2.358) and VLDL (6.666±0.666). Generally, liver function test (ALT and AST) increased significantly in male rats treated with three dose of Methomyl during 28 days. ALT level in G1 was (139.333±5.931), in G2 was (134.500±3.461) and in G3 was (125.166±9.775) when compared to the control group (119.000±2.542). AST level was (41.500±1.431) in G1, (40.666±1.763) in G2 and (50.166±4.860) in G3 when compared with control group (37.450±3.471). On the other hand, the level of ALP was found to be non-significantly increased in three dose Methomyl administration (287.333±22.340, 306.166±16.541 and 309.000±28.183) in G1, G2 and G3 respectively when compared to the control group (160.833±10.581).

Table (2): Effect of Methomyl on the Lipid Profile in Male Rats and Control Group after 28 days.

<table>
<thead>
<tr>
<th>Group</th>
<th>Control (0.5 mg/kg bw)</th>
<th>G1 (1 mg/kg bw)</th>
<th>G2 (1 mg/kg bw)</th>
<th>G3 (2 mg/kg bw)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum Urea (mg/dl)</td>
<td>21.261±0.299</td>
<td>20.350±0.888</td>
<td>18.416±0.808</td>
<td>17.600±0.524</td>
<td>0.096</td>
</tr>
<tr>
<td>Serum Creatinine (mg/dl)</td>
<td>0.540±0.025</td>
<td>0.480±0.022</td>
<td>0.483±0.01</td>
<td>0.545±0.015</td>
<td>0.739</td>
</tr>
<tr>
<td>Cholesterol (mg/dl)</td>
<td>62.000±0.862</td>
<td>65.500±1.335</td>
<td>79.000±1.861</td>
<td>69.833±3.590</td>
<td>0.001</td>
</tr>
<tr>
<td>TG (mg/dl)</td>
<td>34.333±3.382</td>
<td>41.166±4.206</td>
<td>49.833±5.940</td>
<td>43.000±4.618</td>
<td>0.269</td>
</tr>
<tr>
<td>HDL (mg/dl)</td>
<td>28.900±2.487</td>
<td>24.133±0.717</td>
<td>24.567±1.322</td>
<td>20.933±0.581</td>
<td>0.009</td>
</tr>
<tr>
<td>LDL (mg/dl)</td>
<td>26.833±2.358</td>
<td>31.666±1.666</td>
<td>44.266±1.150</td>
<td>38.666±2.691</td>
<td>0.246</td>
</tr>
<tr>
<td>VLDL (mg/dl)</td>
<td>6.666±0.666</td>
<td>8.333±0.881</td>
<td>8.500±1.118</td>
<td>7.833±0.872</td>
<td>0.876</td>
</tr>
<tr>
<td>ALT (U/L)</td>
<td>119.000±2.542</td>
<td>139.333±5.931</td>
<td>134.500±3.461</td>
<td>125.166±9.775</td>
<td>0.009</td>
</tr>
<tr>
<td>AST (U/L)</td>
<td>37.450±3.471</td>
<td>41.500±1.431</td>
<td>40.666±1.763</td>
<td>50.166±4.860</td>
<td>0.000</td>
</tr>
<tr>
<td>ALP (U/L)</td>
<td>160.833±10.581</td>
<td>287.333±22.340</td>
<td>306.166±16.541</td>
<td>309.000±28.183</td>
<td>0.099</td>
</tr>
</tbody>
</table>
The effect of repeated doses of Methomyl insecticide on body weight gain of the male rats was recorded in Table (3) The findings indicated that male rats before treated was (313.00±1.612, 361.33±1.145 and 380.00±2.708) for G1, G2 and G3, after treated with three dose Methomyl for 28 days showed significant increase in their final body weights, in G1 was (363.17±4.743), in G2 was (428.00±2.098) and in G3 was (436.00±6.465) as compared with control group (350.67±9.383).

Table (3): Effect of Methomyl on Body Weight gain of The Male Rats after 28 Days.

<table>
<thead>
<tr>
<th>Body Weight (g)</th>
<th>Control (0.5 mg/kg bw)</th>
<th>G1 (1 mg/kg bw)</th>
<th>G2 (2 mg/kg bw)</th>
<th>G3 (4 mg/kg bw)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Before</td>
<td>289.67±1.687</td>
<td>313.00±1.612</td>
<td>361.33±1.145</td>
<td>380.00±2.708</td>
<td>0.439</td>
</tr>
<tr>
<td>After</td>
<td>350.67±9.383</td>
<td>363.17±4.743</td>
<td>428.00±2.098</td>
<td>436.00±6.465</td>
<td>0.034</td>
</tr>
</tbody>
</table>

Table (4) show the effects of methomyl administration for 28 days on organs weight (liver and kidney), weight of liver increased significantly in all three groups (11.498±0.112, 13.370±0.317 and 13.615±0.722) in G1, G2 and G3 respectively as compared to control group (11.036±0.206). According to kidney weight, the right and left part increase in the weight which were (1.363±0.062) in G1, (1.598±0.080) in G2 and (1.583±0.0187) in G3, but left part was (1.475±0.060) in G1, (1.591±0.106) in G2 and (1.580±0.034) in G3 in comparison with control group for right and left part (1.320±0.062 and 1.420±0.057) respectively.

Table (4): Effect of Methomyl on Organ Weight of the Male Rats and Control Group after 28 Days.

<table>
<thead>
<tr>
<th>Group</th>
<th>Control (0.5 mg/kg bw)</th>
<th>G1 (1 mg/kg bw)</th>
<th>G2 (2 mg/kg bw)</th>
<th>G3 (4 mg/kg bw)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Liver (g)</td>
<td>11.036±0.206</td>
<td>11.498±0.112</td>
<td>13.370±0.317</td>
<td>13.615±0.722</td>
<td>0.000</td>
</tr>
<tr>
<td>Kidney (g)</td>
<td>Right</td>
<td>1.320±0.062</td>
<td>1.363±0.062</td>
<td>1.598±0.080</td>
<td>1.583±0.0187</td>
</tr>
<tr>
<td></td>
<td>Left</td>
<td>1.420±0.057</td>
<td>1.475±0.060</td>
<td>1.591±0.106</td>
<td>1.580±0.034</td>
</tr>
</tbody>
</table>
Histopathological Changes In Liver And Kidney

Normal liver structure appeared in the form of hepatic lobules in which there were centrally located central veins, which were surrounded by hepatocytes arranged in the form of hepatic cords separated from each other by hepatic sinusoids (Fig. 1A). The liver of rats treated with methomyl at dose level 0.5 mg/kg bw, after 28 days of treatment was normal as control (Fig. 1B). However, the liver of rats treated with methomyl at dose level 1 and 2 mg/kg bw, after 28 days of treatment showed degenerative changes in the liver including dilation of central vein, congestion blood vessels, infiltration and vasodilatation (Fig. 1C and 1D).

Figure.1.a. Photomicrograph of normal architecture with normal hepatocytes central vein (cv), hepatic cells (hc) sinusoids(s). Control group. Haematoxylin/eosin staining; 400 x.
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Figure 1.b. Photomicrograph of rats treated with 0.5 mg/kg bw methomyl showing normal histological structure. Normal hepatocytes, central vein (cv), hepatic cells (hc) haematoxylin/eosin staining; 100 x.

Figure 1.c. Photomicrograph of treated rats with 1 mg/kg bw methomyl showing dilation of central vein, infiltration (inf), vasodilatation. Haematoxylin/eosin staining; 4100 x.
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Figure 1.d. Photomicrograph of rats treated with 2 mg/kg bw methomyl showing dilation of central vein, congestion blood vessels (con), advanced infiltration (inf) and vasodilatation. Haematoxylin/eosin staining; 400x.

In normal histologic structure of the kidney, the cortex contains glomerular tufts scattered in between proximal and distal convoluted tubules (Fig. 2A). Kidney of rats treated with Methomyl at dose level 0.5 mg/kg bw, after 28 days of treatment showed no changes (Fig. 2B). However, for rats treated with Methomyl at dose 1 and 2 mg/kg bw, after 28 days of treatment caused destruction of the normal pattern of the renal tissue. These damages were encountered by the presence of shrinkage of glomerular tuft, thickening of parietal layer of Bowman’s capsule, glomerular degeneration as well as cellular infiltration is clearly seen and renal medulla showed dilated collecting tubules stuffed with R.B.Cs. (Fig. 2 C, D and E).
Figure 2.a. Photomicrograph of the kidney from control rat, showing the normal histological structure of renal parenchyma, rounded glomerulus (g), normal kidney tubules (kt) with narrow lumen lined by cuboidal epithelium. Haematoxylin/eosin staining; x100.

Figure 2.b. Photomicrograph of rats treated with 0.5 mg/kg bw methomyl showing normal histological structure. Glomerulus (g), normal kidney tubules (kt) haematoxylin/eosin staining; x100.
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Figure 2.c. Photomicrograph of treated rats with 1mg/kg bw methomyl from renal cortex showing hypertrophy of glomerular tuft (hy), thickening of parietal layer of bowman’s capsule (th) and infiltration (inf). Haematoxylin/eosin staining. 400x.

Figure 2.d. Photomicrograph of rats treated with 2 mg/kg bw methomyl from the renal cortex showing focal area of mononuclear cellular infiltration (inf), hemorrhage in the lumen of renal tubules (he). Haematoxylin/eosin staining; 400x.
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Figure 2.e. Photomicrograph of treated rats with 2 mg/kg bw methomyl from kidney showing shrinkage of glomerular tuft (sh), infiltration (inf) and dilation of the lumen of renal tubules (d). Haematoxylin/eosin staining; 400x.

Discussion

In the present study, the RBC’s count, Hb concentration and PCV were decreased with MCV, MCH and MCHC in methomyl treatment. The anemia observed might have been due to the ability of the methomyl to cause acute hemolysis or it might be due to cause oxidative stress, these changes due to an increase rate of breakdown of red cells and/or the toxic effect of methomyl on bone marrow. (14). The reduction in Hb content may be due to increased rate of breakdown of red cells and/or reduction in the rate of formation of RBC’s. The present result is similar to that of (15) who observed a decrease in RBC’s count, Hb concentration and PCV in rats exposed to methomyl (1.70 mg/kg bw) for 90 day. Creatinine is excreted completely in urine via glomerular filtration, an elevation of its level in the blood is indicator for impaired kidney function (16). When liver cell plasma membrane is damaged, a variety of enzymes normally located in the cytosol are released into the blood stream. Their estimation in the serum...
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is a useful quantitative marker of the extent and type of hepatocellular damage. One of the most sensitive indicators of hepatocyte injury is the release of intracellular enzymes, such as AST, ALT and ALP in the circulation when methomyl used(17). The present results showed a significant increase in the activities of AST, ALT and ALP in the serum of treated rats suggesting that methomyl might cause critical injury to the liver. The increased levels of serum enzymes indicate an improvement of permeability, hepatocyte damage or necrosis (18). AST and ALT are important and critical enzymes in the biological processes, these enzymes are involved in the breakdown of amino acids into α-keto acids which are coured for complete metabolism through the Krebs’s cycle and electron transport chain (19) The histopathological examination results in this study demonstrated that 28 day the oral intake exposure of rat to methomyl at the tested dose 1and 2 mg/kg bw. resulted in degenerative changes in the liver including congestion blood vessels, infiltration, vasodilatation and hemorrhage. The liver recognized as a target organ of the toxic impact regarding its function in biotransformation and excretion of xenobiotic (20). These present results are in agreement with (21) who reported similar histopathological changes including mononuclear cell infiltration, congestion, hydropic degeneration and hepatocellular damage in the liver of male rats treated with dimethoate, endosulfan and carbaryl. Also (22) who found that a 30-day exposure of male rats to dimethoate at doses of 6 and 30 mg/kg caused portal inflammation, centrizonal congestion and focal hepatocyte necrosis in the liver of rats. Nephrotoxicity can result in systemic toxicity causing decreased ability to excrete body wastes, inability to maintain body fluid and electrolyte balance and decreased synthesis of essential hormones (23), the result in this study come in accordance with the results obtained by (24) who reported that exposure of rats to methomyl (2 mg/kg) noticeably affected glomeruli, tubules and interstitium. Glomeruli appeared swollen bowman's spaces. Glomerular swelling was primarily caused by congestion of glomerular capillaries and thickening of glomerular basement membranes (25) and( 26) observed proliferation and swelling of glomerular endothelial cells and tubular degeneration, mononuclear cell infiltration and fibrosis in thiodicarb and carbendazim treated rats, respectively . also the results in this
study agree with the findings of (27) and(28) they reported kidney damage such marked tubular
dilation, hydropic degeneration in tubular lining epithelium, moderate congestion and
hemorrhage in the cortical male Wistar rats exposed some organophosphate pesticides.

**Conclusion**

The results of the present study suggest that Methomyl has the capability to change the
hematological and physiological parameters, in addition dysfunction and tissue degeneration in
liver and kidney of treated rats, and the effect was pronounced with the long time exposure.
Therefore, the application of the Methomyl should be limited to a designed program with
special care in handling to limit or minimize its hazards.

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