Effect of Oral Administration of Formalin on Blood Gases Parameters in Rats

O.K. Al Omari, W.A. Khamas and A. Elbeteiha

Department of Applied Biological Sciences, Faculty of Science, Jordan University of Science and Technology, P.O. Box 3030, Irbid, 22110, Jordan

College of Veterinary Medicine, Western University of Health Sciences, Pomona, CA 91766-1854, USA

Abstract: Forty Sprage-Dawely rats of both sexes (4 to 5 months of age) were used in this experiment. Formalin solution was provided with drinking water ad libitum for 8-12 weeks. Fourteen rats received a daily dose of 80 mg/kg/day, 13 rats received 150 mg kg⁻¹ day while another 13 rats were given tap water. Animals weights, food and water consumption were monitored throughout the experiment. The concentration of the test substance in the solutions were adjusted weekly on the basis of mean body weight and liquid consumption to maintain a target intake of 80 and 150 mg kg⁻¹ day throughout the experiment. Arterial blood after eight weeks were collected from hypnotized rats given Midazolam-3 mg kg⁻¹ intra-peritoneally (Dormicum F. Hoffmann-La Roche, Ltd, Switzerland, 2371081). Animals became partially anesthetized and blood was collected by cardiac puncture from the left ventricle. Standard procedure was followed to store the blood and immediately transported to the laboratory. The samples were analyzed with blood gas analyzer (Omni 4: AVL, Graz, Austria 2493) pH, HCO₃⁻, PO₂, PCO₂, O₂ Saturation, BE (Base Excess). The same procedure and parameters were followed and tested after 12 weeks of exposure with the blood collected from the tip of the tail of anesthetized rats. Besides, blood samples were collected for reticulocytes counts and stained with retic stain (brilliant cresyl blue). Liver and kidney were excised and weighed and the relative organ weights were calculated. Histological examinations were performed on kidney and liver after routinely processed for normal histological procedure. Sections were stained with hematoxylin and eosin stain. Body weight, food and water consumption showed a dose related decrease which were reflected on the general animal health. There were no clear statistical differences between treated and control groups in respect to most blood gases parameters used with the exception of the pH and PCO₂. A significant decrease in pH and PCO₂ in rats ingested 150 mg/kg/day formalin solution for 12 weeks were evident. Liver and kidneys showed no histological changes in treated rats of all groups. It was observed that reticulocytes percentage decreased significantly in a non-dose related manner.

In conclusion, the effect of the exposure to formalin differ from species to species and from system to system according to the availability, activity and amount of the enzyme formaldehyde dehydrogenase present in that specific species or system.

Key words: Formalin, blood gases, rats, parameters, reticulo cytes, hematoxylin

INTRODUCTION

Formaldehyde is the smallest member of the aldehyde series (HCOH). It contains only one carbon atom, described as a colorless flammable gas with a strong pungent suffocating odor. It bears the highest reactivity amongst the aldehydes, polymerizes with itself spontaneously and undergoes many reactions like condensation, oxidation and reduction. It is known by other names like formalin when dissolved in water and paraformaldehyde, which is the solid polymer of the compound (Gerberich et al., 1980).

Formaldehyde is used principally to produce so many products and used in large amounts in industry (Madestau, 1987). It is used as both preservative and fixative of anatomical, histological and pathological specimens. Subjects in such occupations are candidates to chronic exposures of effective concentrations. All of us are continuously exposed to formaldehyde as it results from cigarette, cosmetics and even food preservatives (Feinman, 1988).

Formaldehyde exists in physiological systems in minute concentrations through the metabolism of amino acids and nucleic acids. It is metabolized by the enzyme formaldehyde dehydrogenase, a specific enzyme distributed all over the body with a higher concentration in the liver and blood that converts formaldehyde to formic acid then to CO₂ and water (Restani et al., 1992). This mechanism of elimination face exhaustion when respectful amount are challenged through different routes.
The need for estimation of blood gases parameters would provide useful information on the pathophysiologic toxicity of formaldehyde and would acquire attention to a different potential of the toxin. The aim of the present study is to evaluate the subchronic toxicity of formaldehyde on acid-base balance (if any) and the effect on circulating reticulocytes.

**MATERIALS AND METHODS**

**Animals:** Forty Sprague-Dawley male and female rats age 4 to 5 months were used in this experiment. Rats were provided by the animal house unit at the Faculty of Medicine, Jordan University of Science and Technology. They were maintained in a controlled atmosphere 21±1°C under 12 h light: 12 h darkness schedule. Food and tap water were obtained from the facility and provided ad libitum.

Animals apparently healthy and normal with regards to general body condition. Three rats were selected randomly for gross examination of internal organs after being sacrificed by cervical dislocation under light ether anesthesia. No abnormalities or disorders of any kind were detected in these animals. Animals were divided into groups for different treatment (Table 1).

**Test substance:** Paraformaldehyde, prilled 95% formaldehyde was obtained from Aldrich Chemical Company (Milwaukee, WI 53233, USA). Paraformaldehyde was dissolved in tap water by continuous stirring and heating at 80°C until the solution was clear. The stocks were prepared once a week and stored in the refrigerator at 4°C.

Formalin solution provided to 30 rats ad libitum for 8-12 weeks. Fourteen rats received a daily dose of 80 mg/kg/day, 13 received 150 mg/kg/day and 13 rats were given tap water and served as a control group. Animals were weighed throughout the experiment on a weekly basis. Food and water consumption were measured every day for the first week then for three random days per week for the remaining period.

The concentration of the test substance in the solutions was adjusted weekly on the basis of mean body weight and liquid consumption to maintain a target intake of 80 and 150 mg/kg/day throughout the experiment.

**Arterial blood:** After 8 weeks of formalin ingestion, 14 animals were hypnotized using Midazolam (Dormicum F. Hoffmann-La Roche Ltd, Switzerland, 2371081). Midazolam (3 mg kg⁻¹) was injected intraperitoneally using insulin syringes. After 10-15 min of injection, the animals became partially anesthetized. Using heparinized 5 mL syringes, blood was collected by cardiac puncture from the left ventricle. The needles were then bent, capped, placed in cold icebox (4°C) and immediately transported for analysis with an average time of 30±10 min. The samples were analyzed using blood gas analyzer (Omni 4: AVL, Graz., Austria 2493) for the measurement of pH, bicarbonate (HCO₃⁻), Base Excess (BE), oxygen and carbon dioxide partial pressure (PO₂ and PCO₂) and saturation O₂.

**Capillary blood:** After 12 weeks of formalin oral administration, capillary blood samples were collected from the tip of the tail of 13 rats. Blood samples were collected from unanaesthetized rats from all groups. Blood was collected in heparinized capillary tubes. They were agitated thoroughly to prevent coagulation through ensure mixing with heparin and processed as mentioned above.

**Reticulocyte counts:** Animals ingested 150 mg kg⁻¹ were assigned at the end of the experiment for reticulocytes count. Animals were anaesthetized and blood was collected in syringes by cardiac puncture and placed in vacutainers containing EDTA. Samples were immediately processed as follows: for each sample of 50 μL blood was transferred to eppendorf tube, followed by the addition of equivalent volume of brilliant cresyl blue (retic stain). The mixture was vortexed for few seconds and placed in a 37°C incubator for 15 min. A drop was placed over a glass slide edge and smeared using a cover slip to prepare a blood film. Ten fields from each sample were examined under oil immersion lens. The counts were then expressed as percentages according to the following equation:

\[
\text{Reticulocyte percentage} = \frac{X}{n} \times y \times 100
\]

- **n** = Number of fields examined
- **x** = Total number of reticulocytes for all fields
- **y** = Average number of cells per field

<table>
<thead>
<tr>
<th>Formalin dose mg kg⁻¹</th>
<th>Number of animals</th>
<th>Number of animals 12 weeks exposure</th>
<th>Number of animals for reticulocytes stain</th>
<th>Histological examination of the liver and kidneys</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>6</td>
<td>7</td>
<td>7</td>
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<tr>
<td>80</td>
<td>8</td>
<td>6</td>
<td>6</td>
<td>6</td>
</tr>
<tr>
<td>150</td>
<td>6</td>
<td>7</td>
<td>7</td>
<td>7</td>
</tr>
<tr>
<td>200</td>
<td>20</td>
<td>20</td>
<td>20</td>
<td>20</td>
</tr>
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</table>
Gross anatomy and histology: The same thirteen rats ingested 150 mg kg\(^{-1}\) were sacrificed by cervical dislocation under light ether anesthesia. Liver and kidneys were excised and weighed and the relative organ weights were calculated. Histological examinations using standard technique were performed on kidneys and liver. Staining of all histological tissues were carried out using hematoxylin and eosin stain (Bancroft and Gamble, 2005).

**RESULTS**

General health and gross pathology: General health of rats was estimated through the evaluation of body weights, food and water consumption. Weight gain was significantly reduced in both treatment groups (Fig. 1).

![Graph showing body weight over weeks for different formalin doses](image)

**Fig. 1:** Effect of formalin oral ingestion on body weights of adult male rats. Ingestion of formalin for 12 weeks resulted in a significant decrease in weight gain of rats, which ingested 80 and 150 mg/kg/day in a dose related manner (*p<0.05, **p<0.01*)

The three parameters (body weight, food and water consumption) showed a dose related decrease demonstrating a direct effect of formalin on general health (Table 2). A dose related change was demonstrated in yellow discoloration on the fur that was highly noticeable in the high dose group (150 mg/kg/day) and to a lesser extent in the low group (80 mg/kg/day).

**Blood gas analysis:** After eight weeks, the data obtained by cardiac puncture did not establish any significant statistical differences in the measured parameters among treated and control groups (Table 3). While, after 12 weeks of oral administration, blood samples were collected from the tip of the tail of anaesthetized rats were analyzed for the same parameters mentioned above.

The analysis showed a statistically significant decrease in PCO2 and pH in rats, which ingested 150 mg/kg/day formalin solution (Table 4).

**Reticulocytes counts:** Formalin ingestion for twelve weeks caused a significant increase in the reticulocyte percentage in a non-dose related manner (Fig. 2). The average percentage of reticulocyte in the control group was 1.5 % whereas in the treated groups were 3.3% (*p<0.01*) and 2.9% (*p<0.01*) in the groups ingested 80 and 150 mg/kg/day, respectively.

**Liver and kidney relative weights:** In rats ingested 80 mg/kg/day, a clear significant reduction in relative weights took place in both the liver (*p<0.05*) and the kidney (*p<0.001*). The average relative weights of the liver and kidneys was 3.00 mg ±0.17 and 0.55 mg\(^{-1}\) ±0.05, respectively. On the other hand, liver and kidney relative weight.

**Table 2:** Effect of twelve weeks ingestion of formalin via drinking water on average body weight, water and food consumption in rats exposed to formalin in drinking water for 12 weeks

<table>
<thead>
<tr>
<th>Formalin dose (mg/kg/day)</th>
<th>Body weight (g)</th>
<th>Water consumption ml kg(^{-1})</th>
<th>Food consumption g kg(^{-1})</th>
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</thead>
<tbody>
<tr>
<td>0</td>
<td>391.5±40.2 a</td>
<td>29.38±2.21</td>
<td>28.32±0.49</td>
</tr>
<tr>
<td>80</td>
<td>365.3±38.7</td>
<td>(76.51±5.66)</td>
<td>(72.16±4.27)</td>
</tr>
<tr>
<td>150</td>
<td>345.8±38.2</td>
<td>(50.69±3.13)</td>
<td>(49.73±2.21)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>17.2±0.37**</td>
<td>17.16±2.52**</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(49.88±1.02)**</td>
<td>(50.93±1.39)</td>
</tr>
</tbody>
</table>

*a Results are expressed as mean±standard deviation.\(^1\) Relative water consumption (ml kg\(^{-1}\)).\(^1\) Relative food consumption (g kg\(^{-1}\)).\(^1\) *p<0.05, **p<0.01, ***p<0.001 (student t-test)

**Table 3:** Arterial blood values of rats ingested 0, 80, 150 mg/kg/day formalin solution for eight weeks

<table>
<thead>
<tr>
<th>Formalin Dose (mg/kg/day)</th>
<th>No. of animals</th>
<th>(\text{PO}_2) (mm Hg)</th>
<th>(\text{PCO}_2) (mm Hg)</th>
<th>pH</th>
<th>(\text{HCO}_3^-) (mmol L(^{-1}))</th>
<th>(\text{SO}_2) (%)</th>
<th>BE(^\text{a}) (mmol L(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>6</td>
<td>77.5±4.5(^\text{a})</td>
<td>35.6±2.2</td>
<td>7.43±0.016</td>
<td>23±1.5</td>
<td>95.6±1.0</td>
<td>-0.6±1.3</td>
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<tr>
<td>80</td>
<td>8</td>
<td>75.9±1.12</td>
<td>35.9±2.8</td>
<td>7.44±0.039</td>
<td>24±1.2</td>
<td>95.2±2.1</td>
<td>0.5±1.6</td>
</tr>
<tr>
<td>150</td>
<td>6</td>
<td>80.3±1.08</td>
<td>34.7±4.4</td>
<td>7.42±0.029</td>
<td>22±1.8</td>
<td>95.6±1.8</td>
<td>-1.3±2.5</td>
</tr>
</tbody>
</table>

*a Results are expressed as mean±standard deviation.\(^1\) Base excess
Table 4: Capillary blood gas values of rats ingested 0, 80, 150 mg/kg/day formalin solution for twelve weeks

<table>
<thead>
<tr>
<th>Formalin dose (mg/kg/day)</th>
<th>No. of animals</th>
<th>PO₂ (mm Hg)</th>
<th>PCO₂ (mm Hg)</th>
<th>pH</th>
<th>HCO₃⁻ (mmol L⁻¹)</th>
<th>SO₂ (%)</th>
<th>BE⁺ (mmol L⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>7</td>
<td>30.8±3.5†</td>
<td>55.4±2.9</td>
<td>7.38±0.013</td>
<td>2.1±1.2</td>
<td>57.9±7.1</td>
<td>5.4±0.9</td>
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<tr>
<td>80</td>
<td>6</td>
<td>35.4±10.0</td>
<td>58.3±9.8</td>
<td>7.35±0.045</td>
<td>31.7±3.0</td>
<td>61.7±17.1</td>
<td>4.5±1.6</td>
</tr>
<tr>
<td>150</td>
<td>7</td>
<td>30.6±6.2</td>
<td>61.9±6.3*</td>
<td>7.38±0.198*</td>
<td>33.2±3.3</td>
<td>54.5±12.6</td>
<td>5.6±2.7</td>
</tr>
</tbody>
</table>

*p<0.05, **p<0.005 (student t-test). Results are expressed as mean±standard deviation. † Base excess

Fig. 2: Effect of 12 weeks ingestion of formalin on blood reticulocytes percentage. Data show a significant increase in percentage of reticulocytes in both treated groups compared to the control (p<0.01 for both)

Fig. 3: Effect of 12 weeks oral ingestion of formalin solution on relative liver weights (organ weight (g)/(100g)). Rats that ingested 80 mg kg day showed a significant decrease in liver relative weight (p<0.05)

Fig. 4: Effect of 12 weeks oral ingestion of formalin on relative kidney weight (organ weight (g)/(100g)). Rats that ingested 80 mg kg day showed a significant decrease in kidney relative weight (p<0.05)

of any kind in all groups ingested formalin ingestion with regard to the present experiment.

DISCUSSION

Body weight and water consumption were significantly reduced in a dose-related manner when compared to control. Water consumption is probably due to the high rejection of the formalin solution by rats. These changes were also observed in other study for 4 weeks in rats ingested 125 mg/kg/day (Til et al., 1988). Johannsen et al. (1986) administered formalin to male and female rats and dogs via drinking water for three months. Their results demonstrated that no decrease in food intake, yet water consumption and body weights decreased at 100 and 150 mg kg⁻¹ body weight/day. The differences between rats and dogs in food consumption and feed efficiency are related to some variations in how these two species metabolize formaldehyde.

In a chronic toxicity study, formalin solution (300 mg/kg/day) was administered orally to rats for 24 months. Accelerated death rates, retarded growth, decreased food and water intake were observed (Tobe et al., 1989).

Beall and Ulsamer (1984) suggested that exposure to formaldehyde by various routes including oral administration might be associated with hepatotoxicity in
several animal species. Other studies could not find any hepatotoxic activities in organ weight, histological analysis and liver enzyme activities (Til et al., 1988, 1989; Hurni and Ohder, 1973; Johannsen et al., 1986). In the present study, histology of the liver and kidney were evaluated and the results obtained are in agreement with the previous studies in which no histological changes for both organs were observed in all treatment groups. It seems that formalin action is decreased in the blood and liver to a level that can be tolerated by these organs at the concentration used in this experiment.

Two methods were used in the present study to evaluate the acid-base disturbances caused by formalin ingestion with regard to sampling techniques, arterial (cardiac) and capillary (tail).

Arterial blood showed no clear statistical correlation between treated and control groups in respect to blood gases parameters and formalin ingestion.

Capillary blood samples were collected from the tip of the tail of unanesthetized rats. Data showed significant decrease in blood pH (p<0.005) in rats ingested 150 mg/kg/day and increase in PCO₂ (p<0.05) in the same group. Blood pH of the high dose group indicates a state of acidosis. Unfortunately, data obtained from PCO₂ alone is insufficient to interpret the changes in pH or to identify its nature because PCO₂ is a respiratory component and can only indicate a respiratory abnormal action. Neither of the tests showed a difference in [HCO₃⁻] between control and treated groups which made it difficult to interpret the exact kind of slight acidosis observed in this experiment.

Steave and Haskins (1977) demonstrated that blood samples stored in ice water can maintain pH values, PCO₂ and PO₂ without change for 3.5 and 6 h, respectively. Arterial blood samples were collected after the animals were heptomized but not fully anesthetized to minimize the influence of anesthesia on acid base equilibrium knowing that anesthesia tends to cause ventilatory depression and thus raise the PCO₂ values (Bar-Ilan and Marder, 1980, Brun-Pascaud et al., 1982) (although a slight decrease in pH is expected by heptomization yet to lesser degree). However, both control and treated groups were treated the same way, therefore, changes were most probably due to the formalin. In contrast, samples collected from the tip of the tail over the influence of dilution because the volume of heparin and blood are constant for all samples. Yet, awake rats may vary remarkably with regard to stress and ventilatory rate but the effect of heptomization is eliminated.

Rats have slightly more alkaline blood than humans do (Upton and Morgan, 1975). In the present study, arterial blood samples showed a pH of 7.43 and a capillary blood samples of pH value 7.38. The difference of pH between the two samples is attributed to the low concentration of the HCO₃⁻ in the arterial blood compared to the capillary blood. Moreover, PO₂ and O₂ saturation are much higher in arterial blood than capillary blood while PCO₂ of capillary blood is greater than that in arterial blood sample.

The half-life of formaldehyde in the blood is very short (1.5 min) where it will be converted to formic acid by the enzyme formaldehyde dehydrogenase (Restani et al., 1992). This mechanism of elimination face exhaustion when respectful amount are challenged through different routes. It seems that 80 and 150 mg kg⁻¹ are bearable in rats in respect to blood gases parameters with few exceptions.

In the present experiment, reticulocyte percentage was examined. It was evident that formalin ingestion caused a rise in reticulocyte percentage. This might be attributed to the additive stress efffience of the test substance over the animals, which may affect the capability of the red blood cells to carry enough oxygen and thus results in a premature release of reticulocytes into the blood stream.

It seems that each system and each species has its own threshold for formalin. Concentration of formalin used and duration in rats were tolerable in most cases under current study.

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REFERENCES


