Effects of *Lawsonia inermis* L. (Henna) leaves’ methanolic extract on carbon tetrachloride-induced hepatotoxicity in rats

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ABSTRACT

**Background:** Natural products with therapeutic properties such as plants, minerals, and animal products, for many years, were the main sources of drugs for the treatment of numerous diseases; hence selection of *Lawsonia inermis* L. (Henna) to study its hepatoprotective activity was considered. **Objectives:** This was an attempt to evaluate the hepatoprotective effect of *L. inermis* leaves’ methanolic extract on carbon tetrachloride (CCl₄)-induced hepatotoxicity in rats. **Materials and Methods:** The *L. inermis* leaves’ methanolic extract, which obtained by maceration, was orally administered in doses of 100 mg/kg and 200 mg/kg to the tested animals to assess its effects on serum levels of hepatotoxicity parameters, alanine aminotransferase (ALT), aspartate aminotransferase (AST), alkaline phosphatase (ALP), bilirubin, and total proteins along with histopathological liver sections examination, while silymarin (25 mg/kg), a potent hepatoprotective drug, was used as standard control. **Results:** The two doses of the plant extract showed dose-dependent hepatoprotective effect, as evident by the significant reduction (\(P < 0.05\)) in serum levels of AST, ALT, ALP, and bilirubin along with the improvement in histopathological liver sections compared to CCl₄-only treated animals. **Conclusion:** As experimentally evident, it could be concluded that this plant material could provide a hepatoprotective effect that could be attributed to its antioxidant properties.

KEY WORDS: Carbon tetrachloride hepatotoxicity, hepatoprotective, *Lawsonia inermis* leaves, liver enzymes

INTRODUCTION

Because liver plays important functions in the maintenance, performance, and regulating homeostasis of the body, maintenance of a healthy liver is essential for the overall well-being of an individual [1]. In spite of the tremendous advances in modern medicine, there were few drugs available that offer protection to the liver from damage and help to regenerate hepatic cells [2]. Therefore, it is necessary to search for new drugs for the treatment of liver diseases to supplement/replace the currently used ones. Fortunately, the plant kingdom is a valuable source of new medicinal agents, and it has been reported that approximately 25% of modern medications have been derived from plant materials [3,4]. Hepatoprotective plants contain a variety of chemical constituents such as phenols, coumarins, lignans, essential oil, monoterpenes, carotinoids, glycosides, flavonoids, organic acids, lipids, alkaloids, and xanthenes [3]. However, among numerous plant materials used for liver protection, a considerable number of them lack the scientific prove for these claims [5]. Therefore, it is important to follow systematic research methodology to scientifically evaluate the plant materials that claimed to possess hepatoprotective activity.

*Lawsonia inermis* (Henna) is a shrub or small tree cultivated in many regions as an ornamental and commercial dye crop [6]. It is mostly found in the tropical, sub-tropical, and semi-arid zones of Africa (tropical Savannah and tropical arid zones), south Asia, and north Australia [7]. As reported by Varghese et al., (2010) [8], wide range of chemical constituents have been isolated from Henna which includes napthoquinone derivatives (lawsone which is the chief ingredient and the coloring matter in the leaves), phenolic derivatives, coumarins, xanthones, tannins, flavonoids, aliphatic components, triterpenes, sterols and other chemical constituents such as glucose, gallic acid, amino acids, mannitol, trace elements and minerals.

As a medicinal plant, Henna has been used in folk remedy as astringent, hypotensive, sedative, and against a headache, jaundice, and leprosy [9]. Leaves were also used for skin diseases, venereal diseases, smallpox, and spermatorrhea. Powered seeds
were effective against dysentery and liver disorders. The bark used in a variety conditions, such as burns, jaundice, spleen enlargement, calculus, leprosy, and skin disorders. Root was considered as a potent medicine for gonorrhoea, herpes infection, sore eyes, as an abortifacient, and in the treatment of some nervous disorders [10]. Because, seeds were able to alleviate liver disorders, a question was raised about the protective effects of other plants’ parts.

Therefore, this work was aimed to study the hepatoprotective effect of *L. inermis* leaves’ methanolic extract on carbon tetrachloride (CCl₄)-induced hepatotoxicity in rats as an initial step that can aid further elucidation of the therapeutic potential of this plant product.

**MATERIALS AND METHODS**

**Plants Materials**

Fresh leaves of *L. inermis* were obtained from the local market, Wad Medani, Sudan, which was further identified at the herbarium of the phytochemistry and taxonomy Department, Medicinal and Aromatic Plants Institute, National Center for Research, Khartoum, Sudan, and a voucher specimen was placed for future referencing.

**Extraction of Plants Materials**

The leaves of *L. inermis* were washed with distilled water; shade-dried at room temperature for 72 h, then grinded into powder by glass mortar and pestle. A 300 g of dry powder were extracted by maceration using 1.5 liters of methanol (99%) as a solvent system in conical flasks for 72 h, with intermittent shaking, and then filtered under vacuum using Buchner funnel. The filtrate was allowed to evaporate at room temperature, in dark, for 7 days, and the greenish-black semisolid extract was collected and stored in an amber glass container in a refrigerator until used for biological testing.

**Experimental Animals**

Albino rats weighing about 130-200 g (12-15 weeks old) males and females obtained from the animal house, Faculty of Pharmacy, University of Gezira. They were housed in polycrylic cages and acclimatized to laboratory condition for 14 days before commencement of the experiment (temperature 25 ± 2°C with dark and light cycle 14/10 h). The animals were allowed free access to diet and water. They were divided into five groups as follow:

- **Group I** (normal control, *n* = 5) received distilled water
- **Group II** (negative control, *n* = 5) received CCl₄ only
- **Group III** (standard control, *n* = 5) received silymarin (25 mg/kg)
- **Groups IV and V** (test control, *n* = 10) received *L. inermis* leaves extract 100 mg/kg and 200 mg/kg, respectively.

Water suspensions of silymarin and *L. inermis* extract were daily orally administered to the animal groups (III, IV, and V), respectively. Carbon tetrachloride (CCl₄) was administered every 72 h with intraperitoneal injection in a dose of 2 ml/kg [11] to all rats except the rats of Group I.

This study was ethically approved by the Faculty of Pharmacy, University of Gezira ethical committee, and in compliance with the Faculty of Pharmacy Guiding Principles in the Use of Animals in Toxicology 2006.

**Determination of Hepatoprotective Effect**

Hepatoprotective effect of *L. inermis* extract on CCl₄-induced hepatotoxicity in rats were carried out for 21 days based on the methods described by Sapakal et al., [12] and Azeem et al., [13].

Blood samples were collected at day 7, day 14, and day 21, allowed to clot for 45 min at room temperature, and serum was separated by centrifugation at 3500 rpm for 15 min at 4°C then used for measurement of alanine aminotransferase (ALT), aspartate aminotransferase (AST), alkaline phosphatase (ALP), total bilirubin, and total proteins levels using an auto-analyzer (Cobas Integra 400, Roche Diagnostics, Switzerland).

**Histopathological Studies**

Slices of liver from each animal in all groups were preserved in 10% buffered neutral formalin, the tissues were mounted in the laboratory by embedding paraffin sections, then stained with hemotoxyline and eosin and subjected to histopathological examination at the Histopathology Department - Medical Laboratory - University of Gezira.

**Data Analysis**

The biochemical data were statistically analyzed using paired *t*-test and expressed as mean ± standard error of mean. For comparisons with the CCl₄-only treated group (Group II), differences were considered significant if *P* < 0.05. The percentages of hepatoprotection (H) were calculated as described by Singh et al., [14], using the mean value of each group by the following equation:

\[
H = \left[ 1 - \left( \frac{\text{Group treated with test drug}}{\text{Normal control group}} \right) \right] \times 100
\]

**RESULTS**

The obtained biochemical data showed that administration of CCl₄ to rats significantly elevated the serum levels of ALT and AST compared to the normal group (*P* < 0.01) which indicated acute hepatocellular damage. A significant reduction (*P* < 0.001) in ALT and AST levels were recorded in Silymarin treated group (Group III) compared to CCl₄-alone treated group.
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(Groups II) at day 7 and day 14 of the experimental course. However, at day 21, all treatment groups (Groups III, IV, and V) showed a significant reduction ($P < 0.01$) in ALT and AST levels with the greatest reduction recorded in Group V [Table 1].

The serum levels of ALT, AST, ALP, total bilirubin, and total proteins that obtained from the tested animals at day 21, along with the percentage of hepatoprotection for each biochemical parameter, showed that both plant extract and silymarin decreased the levels of liver enzymes that elevated by CCl₄ administration when compared to CCl₄-only treated group. The two different concentrations that used in this study for the plant extract showed a concentration-dependent effect [Table 2].

In the histopathological analysis, the liver section of Group I (normal control) exhibited normal hepatic cells with distinct hepatic cells, sinusoidal spaces, and central vein [Figure 1], whereas that of CCl₄-only intoxicated animals showed necrosis of hepatic architecture with vacuolization and congestion of sinusoids [Figure 2]. Treatment with silymarin, and the methanolic extract of _L. inermis_ leaves, showed minimal necrosis and regeneration of hepatocytes compared to the CCl₄-only intoxicated group which provided supportive evidence for the biochemical analysis [Figures 3 and 4].

DISCUSSION

In animal model studies, induction of liver injury by CCl₄ is commonly used to evaluate the of hepatoprotective agents, because CCl₄ administration significantly elevated the serum levels of ALT, AST, ALP, and bilirubin [15] that considered as mutual signs for liver injury. CCl₄ bio-transformed by cytochrome P₄₅₀ system in the endoplasmic reticulum to produce trichloromethyl free radical which combined with cellular lipids and proteins in the presence of oxygen form trichloromethyl peroxyl radical, and may attack lipids on the membrane.

**Table 1:** Effects of methanolic extract of _L. inermis_ leaves and Silymarin on serum levels of ALT and AST in CCl₄-induced hepatotoxicity

| Group                  | Day (7)       | Day (14)      | Day (21)      |
|------------------------|---------------|---------------|---------------|
|                        | ALT (U/L)     | AST (U/L)     | ALT (U/L)     | AST (U/L)     | ALT (U/L)     | AST (U/L)     |
| Group I (normal control)| 90±12         | 134±14        | 63±3          | 126±9         | 94±5          | 119±16        |
| Group II (CCl₄ only)   | 3165±190      | 4483±284      | 1826±367      | 2897±81       | 1344±121      | 2259±117      |
| Group III (CCl₄+Silymarin 25 mg/kg) | 791±61 | 1220±115      | 1565±178      | 1985±112      | 214±26        | 629±99****    |
| Group IV (CCl₄+L. inermis 100 mg/kg) | 2536±253      | 3460±216      | 1430±176      | 2800±200      | 691±51        | 1417±45**     |
| Group V (CCl₄+L. inermis 200 mg/kg) | 1874±282      | 3076±168      | 955±65        | 1995±148      | 341±21        | 777±49****    |

**Table 2:** Hepatoprotective percentage of _L. inermis_ leaves methanolic extract and Silymarin on serum levels of ALT, AST, ALP, bilirubin, and total proteins at day 21

| Group                  | ALT (U/L) | AST (U/L) | ALP (U/L) | Bilirubin (mg/dL) | Total proteins (g/dL) |
|------------------------|-----------|-----------|-----------|-------------------|-----------------------|
| Group I (normal control)| 94±5      | 119±16    | 81±5      | 0.19±0.02         | 6.8±0.12              |
| Group II (CCl₄ only)   | 1344±121  | 2259±117  | 383±23    | 1.4±0.08          | 5.4±0.2               |
| Group III (CCl₄+Silymarin 25 mg/kg) | 214±26 (90%)**** | 629±99 (76%)*** | 255±15 (42%)** | 0.3±0.06 (79%)*** | 5.7±0.2 (21)%*      |
| Group IV (CCl₄+L. inermis 100 mg/kg) | 691±51 (52%)*** | 1417±45 (39%)* | 334±14 (16)%* | 0.3±0.07 (79%)*** | 5.6±0.14 (14)%*      |
| Group V (CCl₄+L. inermis 200 mg/kg) | 341±21 (80.2%)**** | 777±49 (69.3%)*** | 311±12 (24)%* | 0.3±0.06 (79%)*** | 5.6±0.12 (14)%*      |

**P value:** **P<0.05, ****P<0.001 versus CCL₄ group, CCL₄: Carbon tetrachloride, _L. inermis_: Lawsonia inermis, ALT: Alanine aminotransferase, AST: Aspartate aminotransferase, ALP: Alkaline phosphatase

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Figure 1: Histopathological liver section of normal control group (H&E, ×10)

Figure 2: Histopathological liver section of carbon tetrachloride-only intoxicated group (H&E, ×10)
of endoplasmic reticulum faster than trichloromethyl free radical [16]. The intraperitoneal route was found to be the best route for CCl₄-induced hepatotoxicity in rats, and the optimum dose was found to be 2 ml/kg (dissolved in an equal volume of olive oil), and this significantly increased the level of serum enzymes, without causing the death of the animals [11].

The serum levels of aminotransferases were markedly elevated by hepatocytes damage due to toxins, drugs, or viruses [17], and estimating the activities of the serum marker enzymes could make a useful quantitative biomarker of the extent and type of hepatocellular damage, and the tendency of these enzymes to return to near normal levels considered as a clear manifestation of antihepatotoxic effects of the administered agent [18].

Hepatoprotective agents exert their action against CCl₄-induced liver injury by impairment of CCl₄-mediated lipid peroxidation, either through decreased production of free radical derivatives or due to the antioxidant activity of the protective agent itself [19]. It has been reported that L. inermis is a rich plant in phenolic compounds such as phenolic acids, flavonoids, tannins, lignin, and others that possess antioxidant, anticarcinogenic, and antimutagenic effects as well as antiproliferative potentials [20].

The obtained results indicated that the tested plant material could protect the liver against CCl₄ toxicity in a dose-dependent manner as evident from the protection provided when compared to the enzyme levels in CCl₄-only treated rats along with histopathological observations. It has been reported that the plant extract affords hepatoprotective activity due to its antioxidant property attributed to the flavonoids content that are effective scavengers of superoxide anions, peroxynitrite, peroxy and hydroxyl radicals [21].

CONCLUSION

It could be concluded that biochemical and histopathological alterations induced by CCl₄ administration were improved under the effect of the plant extract. The concentration of 200 mg/kg was found to be more effective than 100 mg/kg. The present study also suggested that the traditional medicine could have the potential to be transposed successfully in the context of modern medical interventions, but additional researches are necessary to characterize the active constituents and to assess their pharmacological properties.

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