

Evaluation of acute and sub-acute hepatotoxic activity of leaves of *Ehretia laevis* n-hexane extract in mice

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Abstract

Objective: The aim of the present study was to evaluate the acute and subacute hepatotoxicity of *Ehretia laevis* (leaves).

Methods: Toxicity study of *Ehretia laevis* was carried out in Swiss female mice after single ingestion of the extract at different doses (acute toxicity) and daily administration for fifteen days (subacute toxicity).

Results: The results showed that the LD₅₀ of the extract was higher than 4000mg/kg and subacute treatment showed no change in weight of liver however, ALT, ALP, AST bilirubin and globulin values increased significantly whereas albumin and protein decreased. Histopathological studies showed aggregation of cells in parenchyma and in blood vessels vicinity at 1000mg/kg. At 2000mg/kg vacuolation was present, sinusoidal spaces absent, acute cellular swelling and mild degree of cell necrosis was noted. At 4000mg/kg cellular swelling was found to be increased and large number of polymorphic cells was present in blood vessels, showing necrosis. Extract showed dose related toxic effects on hepatocytes.

Conclusions: On the basis of above findings it is concluded that the n- hexane extract of leaves of *Ehretia laevis* has hepatotoxic effect in mice. Further, chronic toxicity studies are needed in other species and with different extract.

Keywords: *ehretia laevis*, n-hexane, mice, hepatotoxicity

Introduction

Ehretia laevis is found world widely in India, China, Australia and Pakistan (Joshi, 2000) [3], locally available in Kashmir and Lahore. Local name of the plant is Chamror and Tamboliya. Its leaves are chewed to cure mouth blisters. For the cure of dysuria leaf powder is taken orally by mixing with sugar and goat milk for 10 days (Jain *et al.*, 2008) [2]. Leaves paste is applied locally in headache, ulcer (Joshi, 2000) [3] and eczema. Flower powder acts as aphrodisiac when taken with milk. Fruit of plant acts as anthelmintic, astringent, demulcent, diuretic, expectorant and also helps in urinary passage affections, lungs and spleen diseases (Joshi, 2000) [3]. Fresh root decoction is used for syphilis and bark of stem for diphtheria. *E. laevis* soaked seeds are ground to make paste, mixed with large cardamomum (*Amomum subulatum*) and taken with milk orally for the cure of jaundice (Sharma *et al.*, 2012) [9]. Seeds act as anthelmintic (Joshi, 2000) [3]. To cure ringworms, powder of kernel is mixed with oil and applied on the affected part. Leaves of the plant contain carbohydrates, proteins, fats, vitamins A, C, E, thiamine and Riboflavin (Torane *et al.*, 2010; Saleem and Naseer, 2014) [11, 8]. Stem and leaves ethanol extract of *E. laevis* shows significant antioxidant activity (Rasika *et al* in 2011) [5]. According to study by Toran & coworkers in 2011, flavonoids and phenolic contents in methanol extract of *E. laevis* leaves shows antioxidant activity *in vitro* by using 2, 2-diphenyl-1-picryl-hydrazyl (DPPH), the study extended with n-hexane extract (Saleem *et al.*, 2015) [7]. In current study n- hexane extract of *Ehretia laevis* leaves are investigated for acute and subacute toxicity as there is no prevention report for evaluation of its toxicity.

Materials and methods

Plant material and extraction

Leaves of the plant used in experimental work were collected

from botanical garden of GC University Lahore, Pakistan. The plant *Ehretia laevis* was compared with the voucher specimen (voucher specimen No. GC Herb. Bot 2288) deposited at Botany Herbarium at G.C. University Lahore. The leaves were washed with tap water; shade dried, powdered and soaked in n-hexane (Perianayagam *et al.*, 2011) for seven days. Extract was prepared by simple maceration process using 5L of n-hexane. Extract was evaporated under reduced pressure using rotary evaporator.

Animals

Thirty Swiss female mice of same age group weighing 25-30g were purchased from National Institute of Health Sciences Islamabad, Pakistan. They were maintained at Animal House of College of Pharmacy G.C. University Faisalabad. They were kept in the cages, had free access to food and water ad libitum and maintained under controlled temperature (23±2°C) with 12h light – dark cycle. The animals were divided into two groups of fifteen mice each. Each group was further divided into control and four treated groups with three mice in each to represent control, 500mg, 1000mg, 2000mg, and 4000mg concentration of the extract in each method. The control group received olive oil, administered by gavage through oral rout for fifteen days.

Selection and preparation of stock solution for toxicity study

The n-hexane extract of the plant was suspended in olive oil to prepare four doses of 500, 1000, 2000 and 4000mg according to animal body weight. Control group was administered with 1ml/kg olive oil. For 500mg /kg of plant extract, by dissolving 50mg of plant extract per one ml of olive oil, the stock solution was prepared and dose was injected as 1ml/100g. For 1000mg/kg dose of each plant extract, 100mg of plant extract

was dissolved per 1ml of olive oil. Similarly 2000 and 4000mg/kg of plant extract were prepared by dissolving 200 and 400mg of plant extract per 1ml of olive oil respectively.

Acute toxicity

The animals were observed for 24h after administering the extract.

Subacute toxicity

Subacute toxicity was observed in all the groups received 500, 1000, 2000 and 4000mg/kg by gavage for 15 days (once a day). The animals were weighed on daily basis. At the end of

experiment animals were sacrificed and blood sample was collected for biochemical analysis. After blood collection, the animals' livers were removed for histopathology. The biochemical parameters evaluated include alkaline phosphatase (ALP), alanine aminotransferase (ALT), aspartate amino transferase (AST), total bilirubin, albumin, protein and globulin.

Statistical analysis

The results were presented as mean ± S.D. and the statistical significance between the groups was analyzed by means of the analysis of variance (one way ANOVA).

Table 1: Effect of treatment with *E. laevis* n-hexane extract on biochemical parameters

Dose mg/kg	Control (n=5)	500mg (n=5)	1000mg (n=5)	2000mg (n=5)	4000mg (n=5)
ALP (U/L)	253.40±2.07	445.80±9.26	426.2±28.5	427±17.04	483.2±19.66
ALT (U/L)	32.60±1.673	58.40±6.50	64.20±0.447	63.60±3.36	68.80±4.09
AST (U/L)	13.80±1.304	26.60±4.16	21.60±5.90	22.20±2.49	27.60±5.50
Bilirubin total	0.640±0.1140	0.8000±0.0000	0.800±0.1225	0.8200±0.0837	0.86±0.1342
Albumin	3.740±0.894	3.0800±0.1095	3.200±0.000	3.200±0.000	3.200±0.000
Protein	6.92±0.0837	6.500±0.300	6.760±0.0548	6.720±0.2049	6.700±0.1225
Globulin	3.12±0.0447	3.580±0.1643	3.340±0.1517	3.420±0.1643	3.6200±0.2168

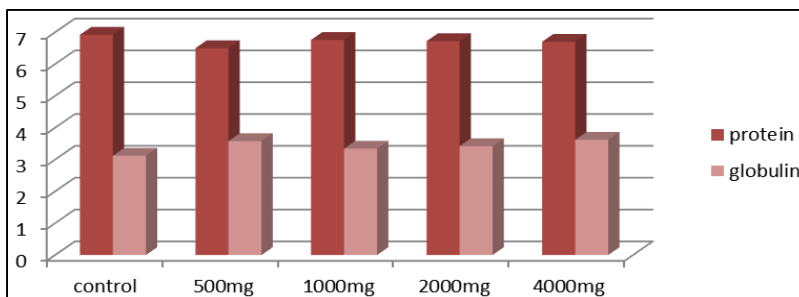


Fig 1: Effects of n-hexane plant extract of *Ehretia laevis* (leaves) on protein and globulin on female mice.

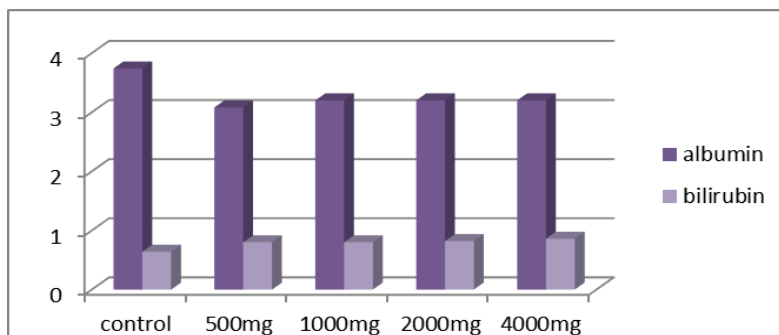


Fig 2: Effects of n-hexane plant extract of *Ehretia laevis* (leaves) on albumin and bilirubin on female mice.

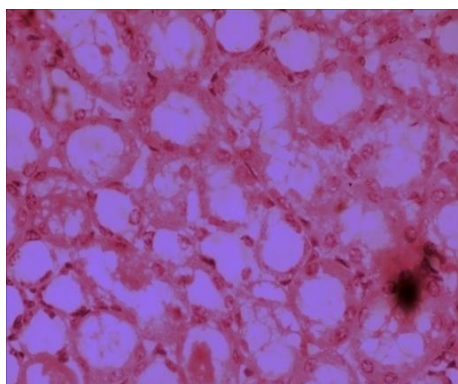


Fig 3: Control group

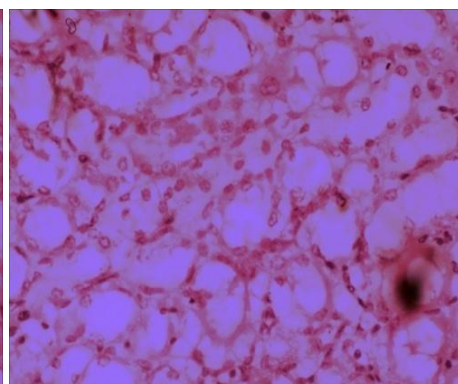


Fig 4: 500mg/kg

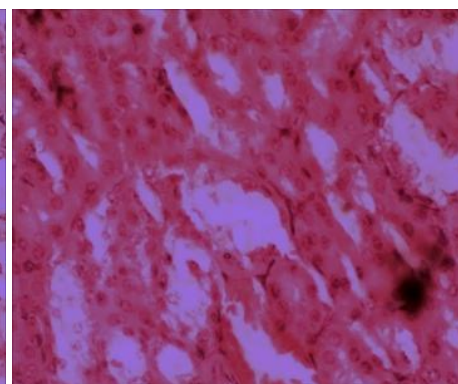


Fig 5: 1000mg/kg

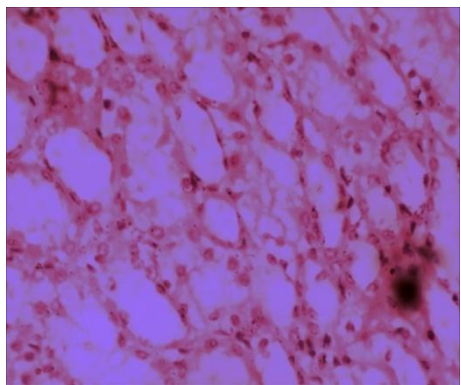


Fig 6: 2000mg/kg

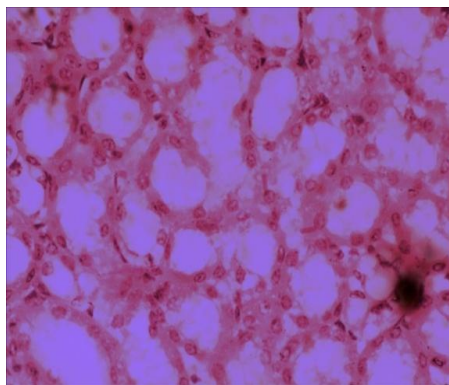


Fig 7: 4000mg/kg

Results and Discussions

Oral administration of n-hexane extract of *E. laevis* in doses from 500-4000mg/Kg did not produce significant changes in breathing, behavior, gastrointestinal effects, and sensory nervous system effects in female albino mice. The effects were observed during the experimental period of (24h). No death occurred in any group during first 24h of experiment. These results showed that there was no adverse effect of *E. laevis* in acute doses, indicating that the median lethal dose (LD50) was higher than 4000mg/kg for mice.

The results showed that n-hexane extract of *E. laevis* is safe in oral administration in acute doses in mice. The multiple administrations with the extract did not decrease water and food consumption (data not shown). The body weight of animals treated with n-hexane extract once a day during 15 days (subacute treatment) did not show any significant change when compared with the control group.

Microscopic analysis of target organ of different animals showed changes in liver structure in comparison with the control group. The extract also showed significant increase in ALP, ALT, AST, bilirubin and globulin values whereas albumin and protein values decreased to some extent (Table 1, Figure 1, 2 and 3). It was also in accordance with the histopathological study, in which control group was compared with the *Ehretia laevis* n- hexane extract. Histopathological studies showed cell aggregates in parenchyma and blood vessels vicinity, at 1000mg/kg. At 2000mg/kg vacuolation was present, sinusoidal spaces absent, acute cellular swelling, less degree of cell necrosis. At 4000mg/kg cellular swelling increased due to sinusoidal spaces, large no. of polymorphic cells were present in blood vessels cells, providing evidence for process of necrosis there. Extracts showed dose related effects on hepatotoxicity. The results were statistically significant.

Conclusion

It was concluded from above mentioned results that plant leaves caused some degree of hepatotoxic injury. Toxicity induced was directly proportional to injected dose i.e. higher the dose greater was the toxicity. However more studies are needed to evaluate chronic toxic effects of *Ehretia laevis*.

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