

Available online on 30.08.2019 at <http://jddtonline.info>

Journal of Drug Delivery and Therapeutics

Open Access to Pharmaceutical and Medical Research

© 2011-18, publisher and licensee JDDT, This is an Open Access article which permits unrestricted non-commercial use, provided the original work is properly cited

Open  Access

Review Article

A Review on Study of Hepatoprotective Activity of *Chenopodium Album* Linnon CCl₄ Induced Hepatotoxicity in Rats

Himanshu Jain, Yogesh Kumar Sharma

Department of Pharmacology, Jaipur College of Pharmacy, Jaipur

ABSTRACT

The hepatoprotective activity of *Chenopodium album* Linn leaves against carbon tetrachloride (CCl₄)-induced hepatotoxicity was investigated. Possibilities of Rat hepatocyte monolayer culture and rats were used as *in vitro* and *in vivo* hepatoprotective screening models also very useful. In the *in vitro* studies, different extracts and fraction we can screened. Silymarin can be used as reference drug. In the *in vivo* studies, hepatotoxicity was induced in wistar rats species give satisfactory results as per reported methods and administering a mixture of CCl₄: olive oil (1:1, 2 ml/kg, s.c.) can be used for the inducible purpose. The extent of hepatotoxicity can be assessed by measuring the serum enzyme levels. So overall parameters consider for the CCl₄ induced hapatotoxicity in rats.

Keywords: Antioxidant; Carbon tetrachloride; *Chenopodium album*; Hepatoprotective

Article Info: Received 02 July 2019; Review Completed 11 Aug 2019; Accepted 22 Aug 2019; Available online 30 Aug 2019



Cite this article as:

Jain H, Sharma YK, A Review on Study of Hepatoprotective Activity of *Chenopodium Album* Linnon CCl₄ Induced Hepatotoxicity in Rats, Journal of Drug Delivery and Therapeutics. 2019; 9(4-A):790-792
<http://dx.doi.org/10.22270/jddt.v9i4-A.3614>

*Address for Correspondence:

Himanshu Jain, Department of Pharmacology, Jaipur College of Pharmacy, Jaipur

1. INTRODUCTION:

1.1 Plant profile: *Chenopodium album*

Chenopodium album is a fast-growing weedy annual plant in the genus *Chenopodium*. Though cultivated in some regions, the plant is elsewhere considered a weed. Common names include lamb's quarters, melde, goosefoot, manure weed, and fat-hen, though the latter two are also applied to other species of the genus *Chenopodium*, for which reason it is often distinguished as white goosefoot. It is sometimes also called pigweed. However, pigweed is also a name for several other plants in the family Amaranthaceae; it is used, for example, for the redroot pigweed (*Amaranthus retroflexus*).[1-3]

It tends to grow upright at first, reaching heights of 10–150 cm (rarely to 3 m), but typically becomes recumbent after flowering (due to the weight of the foliage and seeds) unless supported by other plants. The leaves are alternate and varied in appearance. The first leaves, near the base of the plant, are toothed and roughly diamond-shaped, 3–7 cm long and 3–6 cm broad. The leaves on the upper part of the flowering stems are entire and lanceolate-rhomboid, 1–5 cm long and 0.4–2 cm broad; they are waxy-coated, unwettable and mealy in appearance, with a whitish coat on the underside. The small flowers are radially symmetrical and grow in small cymes on a dense

branched inflorescence 10–40 cm long. Further, the flowers are bisexual and female, with five tepals which are mealy on outer surface, and shortly united at the base.[14] There are five stamens.[4-6]



Figure 1: Plant of *Chenopodium album*

Table1: Plant Description

Kingdom:	Plantae
Clade:	Angiosperms
Clade:	Eudicots
Order:	Caryophyllales
Family:	Amaranthaceae
Genus:	<i>Chenopodium</i>
Species:	<i>C. album</i>
Binomial name	
<i>Chenopodium album</i>	

Chenopodium album Linn, commonly known as 'bathua', 'fathen' or 'lamb's quarters' belongs to family Chenopodiaceae. Due to high nutritional value, leaves are consumed as vegetables in many Asian countries. Traditionally, *C. album* is used as a curative medicine for various diseases including hepatic ailments. Chemically, the presence of phenolics, sterols, vitamins, carotenoids, flavonoids, phytoecdysteroids and minerals has been reported in *C. album* leaves. Many extracts and compounds from *C. album* leaves have been demonstrated to possess hypotensive, anti-inflammatory, antihelminthic, and anticancer activities. Recently the hepatoprotective activity of *C. album* against paracetamol-induced hepatotoxicity has been reported. Our present study was aimed to investigate the protective potential of *C. album* leaves against carbon tetrachloride (CCl₄)-induced hepatotoxicity and oxidative stress.

Liver is the most important organ, which plays a pivotal role in regulating various physiological processes in the body. It is involved in several vital functions, such as metabolism, secretion and storage. It has great capacity to detoxicate toxic substances and synthesize useful principles.

Therefore, damage to the liver inflicted by hepatotoxic agents is of grave consequences. Liver diseases are mainly caused by toxic chemicals, excess consumption of alcohol, infections and autoimmune disorders. Most of the hepatotoxic chemicals damage liver cells mainly by inducing lipid peroxidation and other oxidative damages. In addition, serum levels of many biochemical markers like aspartate transaminase (AST), alanine transaminase (ALT), alkaline phosphatase (ALP) and bilirubin were also elevated. [7-10]

2. EXPERIMENTAL DESIGN AND PROCESS:

2.1 Plant material: The fresh leaves of *C. album* were procured from geographical source place and authenticated by botanist and toxicologist. The fresh leaves were washed with tap water, shade dried and powdered. [11-12]

2.2 species of Animal: Wistar albino rats species (200-250 g) of either sex can be used for the experiments. Animal studies will be reviewed and approved by the Institutional Animal Ethics Committee. [11-12]

2.3 maintenance of animal: The animals were maintained under standard laboratory conditions of temperature (25 ± 2°C) and humidity (55 ± 5%) with 12 h light- dark cycle. [11-12]

3. IN VITRO HEPATOPROTECTIVE ACTIVITY:

Hepatocytes were isolated from rat liver as per the reported method by Jain and Singhai. The isolated hepatocytes were

suspended in William's E medium (pH 7.4) and seeded in collagen pre-coated culture plates at a density of 2 to 3 x 10³ cells/well at 37°C in humidified atmosphere of 5% CO₂ in a CO₂ incubator. After 24 h of culturing, cells were exposed to CCl₄ (2.5 mM) with or without plant extracts/fractions (100 µg/ml) or silymarin (10 µM) and incubated for another 24 h at 37°C in CO₂ incubator. After 24 h incubation, the leakage of alanine transaminase (ALT) and lactate dehydrogenase (LDH) was determined in the culture medium. Acute oral toxicity studies The acute oral toxicity studies were performed following OECD guidelines. [13-14]

4. IN VIVO HEPATOPROTECTIVE ACTIVITY:

The experiment was conducted according to method described previously. Rats were randomly divided into six groups, each consisting of six rats and treated as follows: Group I (normal control): distilled water (1 ml/kg, p.o.) daily for 5 days and olive oil (1 ml/kg, s.c.) on days 2 and 3. Group II (CCl₄ control): distilled water daily for 5 days and CCl₄:olive oil (1:1, 2 ml/kg, s.c.) on days 2 and 3. Group III (positive control): silymarin (50 mg/kg, p.o.) daily for 5 days and CCl₄:olive oil on days 2 and 3, 30 min after administration of silymarin. Groups IV-VI: Extract of plant materials (100, 200 and 400 mg/kg, p.o., respectively) for 5 days and CCl₄:olive oil on days 2 and 3. On the 6th day, under ether anesthesia, blood and liver samples were collected and processed for biochemical estimations. [15]

5. CONCLUSION:

Chenopodium album Linn, commonly known as 'bathua', 'fathen' or 'lamb's quarters' belongs to family Chenopodiaceae. Due to high nutritional value, leaves are consumed as vegetables in many Asian countries. Traditionally, *C. album* is used as a curative medicine for various diseases including hepatic ailments. Chemically, the presence of phenolics, sterols, vitamins, carotenoids, flavonoids, phytoecdysteroids and minerals has been reported in *C. album* leaves. For *in-vitro* The isolated hepatocytes were suspended in William's E medium (pH 7.4) and seeded in collagen pre-coated culture plates at a density of 2 to 3 x 10³ cells/well at 37°C in humidified atmosphere of 5% CO₂ in a CO₂ incubator. After 24 h of culturing, cells were exposed to CCl₄ (2.5 mM) with or without plant extracts/fractions (100 µg/ml) or silymarin (10 µM) and incubated for another 24 h at 37°C in CO₂ incubator. for *in-vivo* Rats were randomly divided into six groups, each consisting of six rats and treated as follows: Group I (normal control): distilled water (1 ml/kg, p.o.) daily for 5 days and olive oil (1 ml/kg, s.c.) on days 2 and 3. Group II (CCl₄ control): distilled water daily for 5 days and CCl₄:olive oil (1:1, 2 ml/kg, s.c.) on days 2 and 3. Group III (positive control): silymarin (50 mg/kg, p.o.) daily for 5 days and CCl₄:olive oil on days 2 and 3, 30 min after administration of silymarin. Groups IV-VI: Extract of plant materials. Protective potential of *C. album* leaves against carbon tetrachloride (CCl₄)-induced hepatotoxicity and oxidative stress.

REFERNCES:

- Asolkar LV, Kakkar KK, Chakre OJ. Glossary of Indian Medicinal Plants with active principles, 1 st part, NISCAIR, New Delhi, India, pp 195-196, 1992.
- Kirtikar KR, Basu BD. Indian Medicinal Plants, 2 nd edition, International Book Distributors, Dehradun, India, pp 2071-2072, 2005.
- Raju M, Varakumar S, Lakshminarayana R, Krishnakantha TP, Baskaran V. Carotenoid composition and vitamin A activity of medicinally important green leafy vegetables. Food Chem 2007; 101:1621-8.

4. Zlatina KN, Paraskev TN, Stefan DN. The genus *Chenopodium*: phytochemistry, ethnopharmacology and pharmacology. *Pharmacog Rev* 2009; 3:280-306.
5. Gohar AA, Elmazar MMA. Isolation of hypotensive flavonoids from *Chenopodium* species growing in Egypt. *Phytother Res* 1997; 11:564-7.
6. Dai Y, Ye WC, Wang ZT, Matsuda H, Kubo M, But PP. Antipruritic and antinociceptive effects of *Chenopodium album* L. in mice. *J Ethnopharmacol* 2002; 81:245-50.
7. Jabbar A, Zamana MA, Iqbal Z, Yaseen M, Shamim A. Anthelmintic activity of *Chenopodium album* (L.) and *Caesalpinia crista* (L.) against trichostrongylid nematodes of sheep. *J Ethnopharmacol* 2007; 114:86-91.
8. Khoobchandani M, Ojeswi BK, Sharma B, Srivastava M. *Chenopodium album* prevents progression of cell growth and enhances cell toxicity in human breast cancer cell lines. *Oxid Med Cell Longe* 2009; 2:160-5.
9. Nigam V, Paarakh PM. Hepatoprotective activity of *Chenopodium album* Linn. against paracetamol induced liver damage. *Pharmacologyonline* 2011; 3:312-28.
10. Kokate CK, Purohit AP, Gokhale SB. *Pharmacognosy*. Nirali Prakashan Publication, Pune, India, pp 109-137, 2005.
11. Jain NK, Singhai AK. Protective effects of *Phyllanthus acidus* (L.) Skeels leaf extracts on acetaminophen and thioacetamide induced hepatic injuries in Wistar rats. *Asian Pac J Trop Med* 2011; 4:470-4.
12. Jain NK, Lodhi S, Jain A, Nahata A, Singhai AK. Protective effects of *Phyllanthus acidus* (L.) Skeels extract on acetaminophen mediated hepatic injury and oxidative stress in Wistar rats. *J Compl Integra Med* 2010; 7. DOI: 10.2202/1553-3840.1439.
13. Robak J, Gryglewski RJ. Flavonoids are scavengers of superoxide anions. *Biochem Pharmacol* 1998; 37:837-41.
14. Jain NK, Singhai AK. Ameliorative effects of *Spinacia oleracea* L. seeds on carbon tetrachloride (CCl₄) - induced hepatotoxicity: In vitro and in vivo studies. *Asian Pac J Trop Biomed* 2012; 1:S232-7.
15. Reitman S, Frankel S. A colorimetric method for the determination of serum glutamic oxaloacetic and glutamic pyruvic transaminases. *Am J Clin Pathol* 1957; 28:53-6.

Journal of Drug Delivery & Therapeutics



JDDT