UTTAR PRADESH JOURNAL OF ZOOLOGY

43(7): 9-14, 2022 ISSN: 0256-971X (P)



Chenopodium album Linn.: A REVIEW ON VARIOUS PHARMACOLOGICAL ACTIVITIES

ANGANA DAS ^{a*¥} AND MRIDUL KR. BORTHAKUR ^{b#}

^a Department of Zoology, Gauhati University, Guwahati-781014, Assam, India. ^b Department of Zoology, B. Borooah College, Guwahati-781007, Assam, India.

AUTHORS' CONTRIBUTIONS

This work was carried out in collaboration between both authors. Both authors read and approved the final manuscript.

Article Information

DOI: 10.56557/UPJOZ/2022/v43i72991

Editor(s):

(1) Dr. Angelo Mark P Walag, University of Science and Technology of Southern Philippines, Philippines.

<u>Reviewers:</u> (1) Isaac John Umaru, Federal University, Nigeria.

(2) Yongwa Gilbert, Cameroon.

(3) Udomdeja Polyium, Rajamangala University of Technology Phra Nakhon, Thailand.

Received: 17 February 2022 Accepted: 26 April 2022 Published: 30 April 2022

Mini-review Article

ABSTRACT

Chenopodium album is widely distributed across the globe and is not only being used in the traditional system of medicine but also in modern medicine. Traditionally, the plant has been used as a hepatoprotective, sedative, diuretic, blood purifier, antiscorbutic laxative and as an anthelmintic against round and hookworms. It is also claimed that the leaves of this plant has a long traditional use in the treatment of digestive, peptic ulcer and hepatic disorder. A comprehensive account on various pharmacological activities like Antioxidant, Antimicrobial, Anti-diabetic, Antipruritic, Antinociceptive, Anti-inflammatory, Anthelmintic and Hepatoprotective activities of *C. album* reported are discussed in this review to explore its immense medicinal properties.

Keywords: Pharmacological activities; *Chenopodium album*; antioxidant; antimicrobial; anti-diabetic; antipruritic; antinociceptive; anti-inflammatory; anthelmintic and hepatoprotective activities.

1. INTRODUCTION

Since the dawn of medicine, diverse groups of people are using medicinal plants to treat various diseases. As the medicinal plants are a reservoir of biologically active compounds with therapeutic properties and minimum side effects, it has drawn the attention of scientists across the globe to use them in the modern medicine system so that the harmful use of chemicals and drugs can be minimized.

Many medicinal plants are used for the treatment of various diseases and one such plant is *Chenopodium album* which is a fast growing annual plant. It falls

^{*}Research Scholar;

[#]Assistant Professor;

^{*}Corresponding author: Email: anganadas94@gmail.com;

under the genus Chenopodium which is distributed in most parts of the world and contains 250 species [1]. About 21 species of this plant are found in India of which some are cultivated to use as a vegetable and a few for the grains obtained from the plant [2]. Naturally, it grows as a weed in the fields of wheat, barley, mustard, gram and other crops [3,4]. The tender shoots are eaten raw and are also cooked as a vegetable. The leaf of this plant is incorporated in various conventional food items to improve the product's nutritional quality and adds variety to the diet [5].

Various studies indicate that C. album is an important dietary element rich in nutrients and antioxidants [6,7]. It is an important source of vitamins like vitamin C and β -carotene which occurs in the young shoots and mature plants of C. album [8]. It is reported that the leaves of this plant have anti-pruritic, antinociceptive activity [9], sperm immobilizing agent cryptomeridiol [10]. and 8-alphaacetoxycyryptomeridiol as growth promoting activity [11]. It is also claimed that C. album contains flavonoid as phenolic amide [12] and has hypotensive activity [13]. Moreover, it is rich in iron [14] and other constituents like saponin [15], cinnamic acid amide [16], alkaloid chinoalbicin [17], apocarotenoids [18], xyloside [19], phenols and ligands [20]. Because of its high nutritional value and medicinal properties, the plant has been used to treat various ailments which occur due to nutritional deficiency. Therefore, for dietary awareness and also to explore the immense medicinal potential of this plant, various pharmacological activities are discussed in this review.

2. PHARMACOLOGICAL ACTIVITIES OF Chenopodium album

2.1 Anti-Oxidant Potential

Medical science has given much emphasis to naturally occurring antioxidants like phenolic acids, polyphenols and flavonoids which inhibits both free radicals and oxidative chain reactions within the tissues and membranes. It is reported that polyphenols are the major plant compounds with antioxidant activity that plays an important role in quenching reactive oxygen species [21]. In 2016, Nowak et al. reported that C. album extracts can be used as a readily accessible source of natural antioxidants and also for food supplement production [22]. In 2017, Lone et al. evaluated the antioxidant activity of C. album by DPPH, riboflavin photo-oxidation, deoxyribose and lipid peroxidation assays. They reported that the extracts exhibited scavenging effect in concentration dependent manner on superoxide anion radicals and hydroxyl radicals [23]. In 2019, Saini et al. studied the antioxidant activity of *C*. *album* by DPPH free radical scavenging activity, total phenolics content and ascorbic acid estimation. Their results justified the nutritional and biological significance of *C. album* [24]. In 2020, Arora et al. carried out a phytochemical screening of *C. album* and reported the presence of flavonoids, tannins, carbohydrates, saponins, proteins and alkaloids [25]. Similar results were also reported by Choudhary et al., [26] and Suleman et al. [27].

2.2 Anti-Microbial Activity

Antibiotics, antifungals, antiprotozoals and antivirals are a group of drugs used to treat microbial infections. Kumar and Kumar [28] studied the antimicrobial activity of methanol and ethyl acetate extracts of C. album against common human pathogens like Klebsicella, P. acne, E. coli, P. aeruginosa, C. albicans and S. cerevisiae. They reported significant antimicrobial activity of methanol extract of C. album against P. acne and S. cerevisiae and mild antimicrobial activity of ethyl acetate extract [28]. To describe the antibacterial activities, Kaur et al. [29] used well plate method where he used three different solvent extracts (methanol, acetone and chloroform) of leaves of C. album against the test organisms Lactobacillus, Bacillus subtilis namely, and Escherichia coli and examined the size of zone of inhibition. The maximum zone of inhibition for 100% concentration was observed as E. coli (19 mm) and Lactobacillus (19 mm) in diameter respectively, while no antibacterial activity was shown against B. subtilis. Compared with standard Amoxicillin, it was found to be 23 mm in diameter for Lactobacillus and 25 mm for both E. coli and B. subtilis in terms of zone of inhibition [29]. Saini et al. [30] mentioned that the antimicrobial effect of methanolic leaf extract of C. album when tested by well diffusion method, proved that the plant possesses considerable inhibitory activity against Escherichia coli, Pseudomonas aeruginosa, Bacillus subtilis, Candida albicans and Candida glabrata [24]. Choudhary et al. [26] also reported antimicrobial activity of C. album during phytochemistry and pharmacological exploration of C. album [26]. Recently, Suleman et al., 2021 also reported the antibacterial activity of C. album against Staphylococcus auerus and S. typhi [27]. A similar report on the antibacterial activity of C. album against Rhizobacteria was also reported by Maria et al., [30].

2.3 Anti-Diabetic Effect

Kant et al. [31] investigated the antidiabetic effect of methanolic extract of *C. album* roots against STZ-induced male Wistar albino rat models. Their results

showed a significant decline in fasting blood glucose levels and a high dose (HD) of C. album extract significantly normalized insulin levels. They further revealed that the methanol extract of C. album roots was effective in normalizing plasma lipid status and decreased cholesterol, triglyceride and LDL levels. Also, a significant decrease in the liver enzymes like SGPT and SGOT was reported [31]. Choudhary et al. [32] evaluated the in vitro and in vivo antidiabetic potential of the flavonoid fraction of C. album and reported potent antidiabetic activity in a dosedependent manner in both in vitro and in vivo diabetic models without any sign of severe toxicity [32]. In a recent review study, Nepal and Chakraborty [33] reported anti-diabetic activity of the plant C. album [33].

2.4 Anti-pruritic and Antinociceptive Activity

To evaluate the crude leaf extract of C. album for central antinociceptive activity in albino mice, Magama and Asita, [34] used Eddy's hot plate test. For antinociception, they used 0, 50, 100 and 150 mg/kg body weight and standard drug Aspirin (150 mg/kg body weight) and found that both Aspirin and the extract at 100 and 150 mg/kg body weight exhibited significant (p<0.05) dose-dependent antinociception compared with the negative control at the seven time intervals; 30, 60, 120, 180, 240, 300, 360 minutes. They found that the maximum antinociception (71.47%) for the 150mg/kg body weight group being at 30 minutes after administration and statistically not different from that of Aspirin (67.44%) at the same time interval. Between 30 and 180 minutes after oral administration of test substances, antinociception due to C. album extract (150 mg/kg body weight) was statistically not different from that due to Aspirin which remained more efficacious than the extract till the end of experiment at 360 minutes; a parallel shift from 240 minutes suggesting a similar mechanism of antinociception [34]. Similar findings on the antinociceptive activity of C. album were also reported by Mushtaq et al., [35]. Choudhary et al. [26] also reported the antinociceptive activity of C. album during phytochemistry and pharmacological exploration of C. album.

2.5 Anti-Inflammatory Activity

Kim et al. studied free radical scavenging activity and enzyme-linked immunosorbent assay (ELISA) experiments on *C. album* extracts to evaluate their anti-oxidative and anti-inflammatory effects. In the free radical (1, 1-diphenyl-2-picrylhydrazyl, DPPH) scavenging activity, EC50 of *C. album* was measured at 0.524 mg/ml. The IL-6 and TNF- α ELISA assay

showed that IL-6 in mouse spleen cells treated with 1 mg/ml of sample decreased the production of IL-6 concentration by 72.30%. In the case of TNF- α , C. album decreased 77.85% of TNF- α production. Their results confirmed antioxidant and anti-inflammatory effects of C. album which can be applied to natural medicine cosmetics having anti-inflammatory effects [36]. Similar findings on the anti-inflammatory activity of C. album were also reported by Mushtaq et al., [35]. Amodeo et al. studied the in vitro antiinflammatory activity of C. album. They reported a significant inhibitory activity on nitric oxide production in lipopolysaccharide-stimulated cells by C. album extract [37]. Choudhary et al. [26] also reported the anti-inflammatory activity of C. album phytochemistry and pharmacological during exploration of C. album.

2.6 Anthelmintic Activity

Anthelmintic or antihelminthic is the substance capable of eliminating parasitic worms (helminths) from the body. It has been reported that helminth stunted infection causes growth, cognitive impairment, anemia and increased susceptibility to other diseases in both human and domestic animals which adds to the economic burden of developing countries [21]. Certain medicinal plants having anthelmintic action has attained great interest for their capability to treat the disease. Lone et al. [23] evaluated the anthelmintic activities of C. album against gastrointestinal nematodes of sheep where they used faecal egg count reduction assay for in vivo study. They reported that the extracts exhibited doseand time- dependent anthelmintic effects on Haemonchus contortus as compared to the standard anthelmintic agent, levamisole. Choudhary et al. [26] also reported the anthelmintic activity of C. album pharmacological phytochemistry and during exploration of C. album. Recently, Choudhary et al., studied the in vitro anthelmintic activity of C. album on earthworms and reported that C. album exhibits two major compounds by LC-MS, i.e., NG and DG, that are mainly accountable for its anthelmintic activity [38].

2.7 Hepatoprotective Activity

Das and Borthakur, 2020 studied the hepatoprotective activity of methanolic leaves extract of *C. album* against paracetamol induced liver damage in albino rats using the biochemical parameters like SGOT, SGPT, ALP, direct bilirubin, total bilirubin and albumin. They reported significant alterations in the increased level of the parameters due to the treatment with the methanolic leaves extract of *C. album* and the results were also supported by histopathological

studies [39]. Some findings on hepatoprotective activity of *C. album* were reported by Parkash and Patel [40]; Aman et al. [41]; Choudhary et al. [26].

3. CONCLUSION

From this review, it can be concluded that the plant C. album possesses important pharmacological activities viz. anti-oxidant, anti-microbial, anti-diabetic, antipruritic. antinociceptive, anti-inflammatory. anthelmintic and hepatoprotective which is of great medicinal value. Also, isolation of pure phytopharmaceuticals and determining their mode of action may lead to the synthesis of novel therapeutic agents. However, there is an evident literature gap regarding the anticancer effect of C. album and phytoanalytical properties of this plant. Therefore, further research on this therapeutically potent herb and its potent toxicity is recommended.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- 1. Risi J, Galwey NW. In advances in applied biology. In: Coaker TH (ed) The *Chenopodium* grains of the Andes: Inca crops for modern agriculture. Academic, Landon. 1984;145-216.
- Yadav N, Vasudeva N, Singh HS, Sharma SK. Medicinal properties of genus *Chenopodium* Linn. Natural Product Radiance. 2007;6:131-134.
- Khurana SC, Malik YS, Pandita ML. Herbicidal control of weeds in potato C.V. kufribadshah. Pesticides. 1986;20:55– 56.
- 4. Bhattacharjee SK. Handbook of medicinal plant. Pointer publishers: Jaipur, 3rd Edition. 2001;01-02.
- Singh L, Yadav N, Kumar AR, Gupta AK, Chacko J, Parvin K, Tripathi U. Preparation of value added products from dehydrated bathua leaves (*Chenopodium album* L.). Natural Product Radiance. 2007;6:06-10.
- Afolayan J, Jimoh FO. Nutritional quality of some wild leafy vegetables in South Africa. International Journal of Food Sciences and Nutrition. 2009;60(5):424-431.
- Hussain J, Khan AL, Rehman NU, Hamayun M, Shah T, Nisar M. Proximate and nutrient analysis of selected vegetable species: A case study of Karak region, Pakistan. African

Journal of Biotechnology. 2009;8(12):2725-2729.

- 8. Gaaza BM.. Niume C. Goduka NI. Nutritional George G. assessment of Chenopodium album L. (Imbikicane) young shoots and mature plant-leaves consumed in the Eastern Cape Province of South Africa. Proceedings International of Chemical, Biological and Environmental Engineering. 2013;53(19):97-102.
- 9. Dai Y, Ye WC, Wang ZT, Matsuda H, Kubo M, But PPH. Antipruritic and antinociceptive effects of *Chenopodium album* L. in mice. Journal of Ethnopharmacology. 2002;81:245-250.
- Kumar R, Mishra AK, Dubey NK, Tripathi YB. Evaluation of *Chenopodium ambrosioides* oil as a potential source of antifungal, antiaflatoxigenic and antioxidant activity. International Journal of Food Microbiology. 2006;115:159-164.
- 11. Pandey S, Gupta RK. Screening of nutritional, phytochemical, antioxidant and antibacterial activity of *Chenopodium album* (Bathua). Journal of Pharmacognosy and Phytochemistry. 2014;3(3):01-09.
- Horio T, Yoshida K, Kikuchi H, Kawabata J, Mizutani J. A phenolic amide from roots of *Chenopodium album*. Phytochemistry. 1993; 33:807-808.
- Gohar AA, Elmazar MMA. Isolation of hypotensive flavonoids from *Chenopodium* species growing in Egypt. Phytotherapy Research. 1997;11:564-567.
- 14. Yadav SK, Sehgal S. *In vitro* and *in vivo* availability of iron from Bathua (*Chenopodium album*) and spinach (*Spinacia oleracea*) leaves. Journal of Food Science and Technology. 2002;39:42-46.
- 15. Lavaud C, Voutquenne L, Bal P, Pouny I. Saponins from *Chenopodium album*. Fitoterapia. 2000;71:338-340.
- Cutillo F, D'Abrosca B, Dellagreca M, Marino CD, Golino A, Previtera L, Zarrellia A. Cinnamic acid amides from *Chenopodium album*: Effects on seeds germination and plant growth. Phytochemistry. 2003;64:1381-1387.
- Cutillo F, D'Abrosca B, DellaGreca M, Zarrelli A. Chenoalbicin, a novel cinnamic acid amide alkaloid from *Chenopodium album*. Chemistry and Biodiversity. 2004;1:1579-1583.
- DellaGreca M, DiMarino C, Zarrelli A, D'Abrosca B. Isolation and phytotoxicity of apocarotenoids from *Chenopodium album*. Journal of Natural Products. 2004;67:1492-1495.

- 19. DellaGreca M, Previtera L, Zarrelli A. A new xyloside from *Chenopodium album*. Natural Product Research. 2005;19:87-90.
- Cutillo F, DellaGreca M, Gionti M, Previtera L, Zarrelli A. Phenols and lignans from *Chenopodium album*. Phytochemical Analysis. 2006;17;344-349.
- 21. Poonia A, Upadhayay A. *Chenopodium album* Linn: review of nutritive value and biological properties. Journal of Food Science and Technology. 2015;52(7):3977–3985.
- Nowak R, Szewczyk K, Dziki UG, Rzymowska J, Komsta L. Antioxidative and cytotoxic potential of some *Chenopodium* L. species growing in Poland. Saudi Journal of Biological Sciences. 2016;23:15-23.
- 23. Lone BA, Chishti MZ, Bhat FA, Tak H, Bandh SA, Khan A. Evaluation of anthelmintic, antimicrobial and antioxidant activity of *Chenopodium album*. Tropical Animal Health and Production. 2017;49:1597-1605.
- 24. Saini R, Kumar D, Mittal A. Antimicrobial and phytochemical potential of *Chenopodium album* Linn. International Journal of Scientific and Technology Research. 2019;8(7):877-880.
- Arora SK, Itankar PR, Yende SR. Phytochemical screening and TLC studies of different extracts of *Chenopodium album*. Journal of Ayurvedic and Herbal Medicine. 2020;6(1):15-20.
- 26. Choudhary N, Prabhu KS, Prasad SB, Singh A, Agarhari UC, Suttee A. Phytochemistry and pharmacological exploration of *Chenopodium album*: Current and future perspectives. Research Journal of Pharmacy and Technology. 2020;13(8):3933-3940.
- 27. Suleman M, Faiz AUH, Abbas FI. Antibacterial, antiparasitic and phytochemical activities of *Chenopodium album* (Bathua) plant extract. Bangladesh Journal of Botany. 2021;50(2):417-421.
- 28. Kumar P, Kumar S. Antimicrobial evaluation and physicochemical study of *Chenopodium album* against some common Human Pathogens. International Journal of ChemTech Research. 2017;10(9):852-858.
- 29. Kaur M, Sharma S, Garg S, Arora M. Study of antibacterial activity of *Chenopodium album* leaves extract. International Journal of Pharmacognosy and Phytochemical Research. 2018;10(1):01-04.
- 30. Maria H, Begum H, Shumail H, Akhtar H, Pervaiz A, Ayaz Y, et al.. Antibacterial activity of *Chenopodium botrys* L. and *Chenopodium*

album L. against growth promoting Rhizobacteria. Pure and Applied Biology. 2021;11(2):586-591.

- 31. Kant S, Dua JS, Lather V. Pharmacological evaluation of antidiabetic and antihyperlipidemic activity of *Chenopodium album* root extract in male Wistar albino rat models. International Journal of Green Pharmacy. 2018;12(2):115-122.
- 32. Choudhary N, Prabhakar PK, Khatik GL, Chamakuri SR, Tewari D, Suttee A. Evaluation of acute toxicity, *in-vitro*, *in-vivo* antidiabetic potential of the flavonoid fraction of the plant *Chenopodium album* L. Pharmacognosy Journal. 2021;13(3):765-779.
- Nepal A, Chakraborty M. An overview on medicinal plants of Sikkim Himalayas region with emphasis on antidiabetic: A review. Journal of Pharmacognosy and Phytochemistry. 2021;10(4):215-217.
- 34. Magama S, Asita AO. Evaluation of *Chenopodium album* Linn. crude methanolic leaf extract for central antinociceptive activity in albino mice using the hot plate test. International Journal of Sciences. 2017;6:36-44.
- 35. Mushtaq A, Rashid S, Jamil M, Anwar R, Khawaja NR. Anti-nociceptive and antiinflammatory activity of *Trapa bispinosa*, *Chenopodium album* and *Cuscuta reflexa*. International Journal of Biology, Pharmacy and Allied Sciences. 2017;6(4):608-622.
- 36. Kim SA, Choi SC, Youn YH, Ko CI, Ha YS, Lee I.A. Antioxidant and Anti-inflammatory effects of *Dioscorea japonica* and *Chenopodium album*. Journal of Society of Cosmetic Scientists of Korea. 2017;43(4):337-347.
- Amodeo V, Marrelli M, Pontieri V, Cassano R, Trombino S, Conforti F, Statti G. *Chenopodium album* L. and *Sisymbrium officinale* (L.) Scop.: Phytochemical content and *in vitro* antioxidant and antiinflammatory potential. Plants. 2019;8(505): 01-14.
- 38. Choudhary N, Khatik GL, Choudhary S, Singh G, Suttee A. *In* vitro anthelmintic activity of Chenopodium album and in-silico prediction of mechanistic role on *Eisenia foetida*. Heliyon. 2021;7:01-08.
- 39. Das A, Borthakur MK. Hepatoprotective activity of Chenopodium album Linn. against paracetamol induced liver damage in albino rats. International Journal of Pharmaceutical Sciences and Research. 2020;11(11); 5605-5610.

- 40. Parkash J, Patel KR. Hepatoprotective activity of *Chenopodium album* leaves extract in CCl₄ induced hepatotoxicity in rats. Journal of Drug Delivery and Therapeutics. 2015;5(2):88-93.
- 41. Aman S, Mazumder A, Gupta UK, Nayak A. Pharmacological activities of *Chenopodium album* Linn.- A review. World Journal of Pharmaceutical Research. 2016;5(10):361-371.

© Copyright MB International Media and Publishing House. All rights reserved.