



CHENOPODIUM ALBUM: A MIRACULOUS TREASURE OF THERAPEUTIC SPECULARITIES

Neelanchal Trivedi*, Bhuvnesh Singh

Teerthanker Mahaveer College of Pharmacy, Teerthanker Mahaveer University, Moradabad, U.P., India

*Corresponding author: neelanchal.trivedi@yahoo.co.in

ABSTRACT

Consciousness regarding medicinal potential of plants is emerging since thousands of years back. But yet it can be concluded that the plant sources remain since the existence of human kind to treat their abnormal physical conditions. It has also been established that excursions, accepted by cultural attributers of aboriginal civilizations, gained appreciable information about the consumption of plants for their therapeutic potentials. *Chenopodium album* is great wellspring of practical supplements and has restorative properties. It can be joined in various expelled sustenance items to make them more nutritious, more advantageous and in addition shopper arranged. The expansion of its leaves to expelled items can upgrade the synthetic and wholesome parameters and can enhance expelled items as practical nourishments. The plants likewise have high natural exercises thus might be of awesome therapeutic esteem. Business abuse of *Chenopodium album* in numerous districts of the world is still a long way from reality. The dynamic constituents can be disengaged and additionally assessed for the advancement of helpful medications. Their cancer prevention agent and antibacterial exercises additionally loan trustworthiness to the natural estimation of this plant.

Keywords: *Chenopodium album*, Herbal Medicine, Antioxidant, Antimicrobial

1. INTRODUCTION

Consciousness regarding medicinal potential of plants is emerging since thousands of years back. But yet it can be concluded that the plant sources remain since the existence of human kind to treat their abnormal physical conditions. It has also been established that excursions, accepted by cultural attributers of aboriginal civilizations, gained appreciable information about the consumption of plants for their therapeutic potentials [1].

Such use of the plants as a medicinal moiety by traditional people placed the foundation for development of today's medicinal system. There are so many companies even in developed part of globe who focused on traditional medicinal system for their formulation development. Medicines from herbal sources are setting trends now a day all over the world. An interesting fact that within USA almost 26% amongst all prescription in the public pharmacies contained the extracts from plant's origin and about 65% of global population remains reliant on traditional system of medicine for their healthcare demand. [2]

Approximately about 8000 medicinal plants are traditionally used alone in India against numerous diseases whereas Korea, China, Malaysia, Japan and few other south-east Asian countries are, in addition, leading in the world in case of consumption of herbal medication.

Nevertheless, there is still the lack of appropriate scientific knowledge on these herbal sources of medicines, which are cultivated as well as utilized in various ailments traditionally by native peoples. This is in any case, rehearsed with no specialized information and thus there is a need to embrace a

logical approach for saving this legacy system of medicine adopted from the beginning of humankind. [3]

To adapt abiotic worries under such conditions, plants are known to blend auxiliary metabolites which can be a rich wellspring of therapeutically vital mixes, for example, flavonoids, terpenoids, tannins, alkaloids and so on. Besides, these plants are wellspring of sustenance, feed, solution, biofuels, consumable oil and so forth in this district and they are ended up being developed as trade trims out numerous parts of the world. In spite of these qualifications, these plants had been generally under investigated for misusing their potential. [4]

Chenopodium album within family Chenopodiaceae is usually called "Bathua" is extensively dispersed in sub-tropical & tropical regions within India. The plant is habitually found in Uttar-Pradesh, Madhya-Pradesh, Himachal-Pradesh, Gujarat, Maharashtra, Haryana, Karnataka, Sikkim, Rajasthan, Jammu and Kashmir & West Bengal. The leaves might be taken as a mixture or the decoction as purgative & anthelmintic. It has likewise been suggested in the therapy of hepatic problems and the enlargement of spleen. The very fine powdered leaves are utilized in form of anthelmintics. It is additionally utilized in stomach problems, eye illness, throat inconveniences, diseases of the blood, heart as well as of spleen and nausea & vomiting. The object of present study was to assess overall impact of *Chenopodium album*. [5]

Consequently, this study has been comprised to assemble all accessible data on the healing approaches of *Chenopodium album* in distinctive pathological conditions with reasonable evidences towards plant's competence.

2. PLANT PROFILE



2.1. Scientific classification: [6]

Kingdom	:	Plantae
Clade	:	Angiosperm
Clade	:	Eudicots
Order	:	Caryophyllales
Genus	:	Chenopodium
Family	:	Amaranthaceae
Species	:	C. album

Common names [7]

Hindi	:	“Bathua”
Sanskrit	:	Vastukah;
Oriya	:	Bathua;
Kannada	:	Kaduoma;Konkani: Chakvit;
Bengali	:	Chandanbethu;
Tamil	:	Paruppukkirai;
Malayalam	:	Vastuccira;
Telugu	:	Pappukura;
Arabic	:	Thanb Alkalb, Rejil Alwaz, Atrah;
Chinese	:	Li;
Unani	:	Bathuaa, Baathu.
Spanish	:	Ceniglo blanco;
English	:	Common goose foot, Lambsquarters, Lamb's-quarters, Fat Hen, Marvel mothi; Pigweed;
Fijian:		Marvel lahan,
French	:	All-good, Muck-weed; Lamb's quarter, Pigweed, Fat hen, Chenopode sauvage (French)
Italian	:	Farinaccio;
Japanese	:	Akaza, Iwa-akaza, Shiroza,
Greek name	:	In greek Chenopodium meaning (goose) & (foot) referring to the shape of leaves of some species.
Latin	:	In Latin, name album means white & alludes to the waxy covering over the plant.

2.2. History and Distribution

It is local of Europe and Western Asia, bathua is an antiquated plant, as per the book (Food in China), bathua has been a nourishment wellspring of a few old civic establishments: it was likely developed in Neolithic Europe (7,000-1700 BC), and was additionally found in China around fifth century AD. Most botanists concur that its inceptions are in reality in Europe, and proof backings the claim that seeker gatherers ate bathua all through the Bronze age and Iron age. Various intriguing accounts portray bathua's surprising history: For instance, Neolithic draftsmen found bathua seeds in early Britain's earthen pots. Scandinavia's Tollund Man, an embalmed carcass of a man thought to live around fourth century BCE, had sheep squatters seeds found in his stomach at his execution site in Danish lowland. Numerous different records specify it as a wellspring of nourishment for the early Vikings, and Peter Kalm's 1749 compositions depict the manners by which Scandinavians heat up the greens in meat-injected water. Indeed, even Napoleon Bonaparte depended on bathua seeds to bolster his troops amid lean circumstances. Inquisitively, archeological leftovers uncover that North American Blackfoot Indian clans were utilizing the weed in the mid-1600s.

However, now, it is one of the most widely distributed species of weeds in the world, especially the temperate zones. [8]

2.3. Morphology

Chenopodium album commonly named as Bathua or Goosefoot, is herbaceous, 0.3-3.5m in height, straight or climbing, mealy or reddish- green, without having any odor.

Stems: Stems once in a while slim, calculated, frequently striped green, red or purple.

Leaves: Leaves are exceptionally factor fit as a fiddle, coming to in developed plants here and there 15cm long, oval, rhombic, deltoid or lanceolated uncaring or intense whole, toothed or sporadically lobulated; petioles-long, slim.

Flowers: Blossoms are in bunches shaping unpredictable or careless panicle regularly coarse spikes, which in developed structures move toward becoming thyrsoid. Sepals-1.5-2mm long, elliptical lanceolated, keeled, shutting over the meagerly membranous utricle.

Seeds: 1.5 mm distance across, orbicular, compacted, with an intense edge, smooth, sparkling, developing life totally annular. The youthful plant of not more than 20cm is quite regarded as a potherb. [9]

2.4. Chemical constituents

Leaves are rich in mineral oil, especially in potash salts, a lot of albuminoids, vitamin C and different mixes are nitrogen. Examination of the leaves gave dampness, 89.6; protein, 3.7; fat 0.4; fiber, 0.8; different starches, 2.9; and minerals, 2.6g;

calcium, 150; phosphorus, 80; press, 4.2 thiamine, 0.01; riboflavin, 0.14; niacine, 0.6; vitamin C, 35mg, carotene 1,740 µg; and vitality, 30kcal/100g; zinc, 24.0; iodine, 0.98; fluorine, 6.3(dry premise); and vitamin K, 250ppm. Betalain alkaloids, phenolic acids in organic products, betain and oxalic corrosive in leaves, oleanolic corrosive and sitosterol in blossoms, furanocoumarins w5x and saponins from the seeds. A phenolic amide has been segregated from the underlying foundations of *C. album*. Its structure was resolved as N-transferuloyl-4-0-methyl-dopamine by spectroscopic proof and substance combination. [10]

2.5. Uses

2.5.1. Traditional uses

The plant was additionally utilized customarily as, anthelmintic against round-and hookworms, antiscorbutic, for treatment of stomach torment, eye ailment, throat inconveniences and cardiovascular issue. Bubbled delicate shoot is utilized as a part of clogging. Fine powder of *Chenopodium album* Linn. Leaves were tidied to partner aggravation & leave's juice was utilized in the treatment of consumes. Decoction of aeronautical fragments blended with liquor was scoured on the body part influenced by joint inflammation and ailment. [11]

2.5.2. Medicinal uses:

In India, the plant is utilized as a purgative, diuretic, narcotic & imbue of the plant is utilized treating the stiffness. It was likewise utilized as an anti-diarrhoeal, anti-phlogistic & antirheumatic, preventative, cardiotoxic, odontalgic, blood purifier & antiscorbutic, stomach related, carminative, love potion, for the treatment of dyspepsia, fart, strangury, original shortcoming, pharyngopathy, splenopathy, hemorrhoids, ophthalmopathy, cardiovascular turmoil, hepatic issue, spleen broadening, biliousness, intestinal ulcers, and general debility. [12].

3. PHARMACOLOGICAL POTENTIAL

3.1. Antioxidant effect

Exhaustive literature revealed the anti-oxidant potential measured and stated as percent inhibition comparative to the control adopting β-carotene bleaching technique. The aqueous & ethanolic extracts of the plant *Chenopodium album* showed the result as 64.5 & 60.5% respectively. The extracts were also subjected further for DPPH radical scavenging as well as BHT scavenging assays and results were significant to those of ascorbic acid. The defensive potential of ethanolic extract was studied on both human & yeast mononuclear leukocytes' genomic DNA compared to oxidative stress. Ethanolic leaf extract of *Chenopodium album* showed marked protection in the DNA fragment of both yeast & human's mononuclear leukocytes when compared to the toxic effect of H₂O₂. [13]

3.2. Antimicrobial activity

Anthelmintic, antimicrobial & insecticidal properties of various extracts of the plant instigated diverged inhibition of used bacterial strains. The antimicrobial efficacy of leaf extract in ethanol of *Chenopodium album* was measured against both gram-positive & gram-negative microorganisms. Bacterial eradicating potential was evaluated against "*Bacillus subtilis*" microorganism with 13.5 mm of ZOI. In-vitro antimicrobial efficacy of the ethanolic & methanolic leave & flower extracts was performed against four strains i.e. *Escherichia coli* & *Pseudomonas aeruginosa* with *Bacillus cereus* & *Staphylococcus aureus*. [14]

In another investigation the aqueous & methanol leaf extracts of *Chenopodium album* were evaluated against i.e. *Salmonella typhimurium*, *Escherichia coli*, *Staphylococcus aureus*, *Proteus vulgaris* & *Pseudomonas aeruginosa*. The result unveiled noteworthy antimicrobial potential against all strains of used bacteria. Meanwhile, the aqueous extract achieved robust antibacterial efficacy against *Staphylococcus aureus* with zone of inhibition of 25 mm whereas slightest antibacterial efficacy was revealed against *Salmonella typhimurium* with zone of inhibition of 17.75 mm. Conversely, methanol leaf extract exhibited significant potential of antimicrobial potential against all included strains of microorganisms. The most relevant potential was calculated with 28.30 mm of ZOI (zone of inhibition) against *Pseudomonas aeruginosa*, though, the bottommost antibacterial efficacy of the plant was detected against *Salmonella typhimurium* with zone of inhibition of 14.00 mm. [15]

However, Amjad and Alizad said that the blooms & leaves of ethanolic & methanolic fractions of *Chenopodium album* don't have any action against the tried bacterial strains. Antifungal action of methanol and n-hexane leaf, stem, root and inflorescence concentrate of *Chenopodium* collection (1, 2, 3 and 4% w/v) was explored against *Macrophomina phaseolina*, a dirt borne contagious plant pathogen that has a wide host run and wide topographical conveyance. The n-hexane concentrates of *Chenopodium* collection lessened parasitic biomass by 60-94%. Two proteins, CAP-I and CAP-II filtered from the leaves of *Chenopodium* collection actuated fundamental protection against tobacco mosaic infection (TMV) and sunnhemp rosette infection (SRV) in both extremely touchy and in addition foundational has. Both CAP-I and CAP-II caused in vitro corruption of TMV RNA. It is recommended that the CAP-I and - II are multi-utilitarian and might act at various levels to guarantee most extreme conceivable hindrance of viral contamination. [16]

The anthelmintic movement of the plant (50, 25 and 12.5 mg/ml) was recorded against grown-up Indian night crawler, *Pheretima posthuma*. In vitro anthelmintic movement of unrefined watery methanolic remove (AME) of *Chenopodium* collection entire plant was considered utilizing full grown *Haemonchus contortus* and their eggs in grown-up motility test and egg bring forth test individually. In vivo anthelmintic

action was assessed in sheep normally tainted with blended types of gastrointestinal nematodes by controlling unrefined powder (CP) and AME in expanding dosages (1.0-3.0 g/kg). Concentrates displayed dosage and time-subordinate anthelmintic impacts by causing mortality of worms and restraint of egg bring forth. LD50 for *Chenopodium* collection was observed to be 0.449 mg/ml in egg incubate test. In vivo, greatest lessening in eggs per gram (EPG) of excrement was recorded as 82.2% at 3.0 g/kg of *Chenopodium* collection AME. Insecticidal impact was applied by the oil ether, carbon tetrachloride and methanol concentrate of *Chenopodium* collection against jungle fever vector, *Anopheles stephensi* Liston. It impacted the early life cycle of *Anopheles stephensi* by lessening the level of bring forth, larval, pupal and grown-up development and furthermore extending the larval and pupal periods. The development file was likewise diminished essentially. [17]

The organic impact of polar and non-polar optional metabolites from the aeronautical parts (leaves and inflorescences) of *Chenopodium* collection against *Oryzaephilus surinamensis* was examined. The outcomes demonstrated that the fluid concentrate of C album was viable with low rate survival of grown-up and larval stages. [18]

3.3. Anti-inflammatory and analgesic effects

The topical anti-inflammatory action for *Chenopodium album* oil (5-0.625 mg) was assessed by hindrance of the 12-O-tetradecanoylphorbol-13-acetic acid derivation (TPA) prompted the edema in ear of mice. The outcome uncovered that the calming activity of the oil is fixation subordinate, the rate diminishment in the ear-edema increments with increment in grouping of the oil. In any case, the oil produced huge diminishment ($p < 0.05$) in edema with all focuses aside from at 0.625 mg [30]. The ethanolic extricate from the products of *Chenopodium* collection, 100-400 mg/kg orally, caused measurement conditionally hindrance of scratching conduct initiated by 5-HT (10 miniaturized scale g for every mouse, sc) or compound 48/80 (50 smaller scale g for every mouse, sc). In any case, it neglected to influence rear paw edema prompted by serotonin or compound 48/80 in mice at measurements of 100 and 200 mg/kg and just demonstrated a moderately feeble restraint on edema at a higher dosage higher than 400 mg/kg. [19]

The part of "NF kappa B (NFkB)" in the antiarthritic capability of concentrates of airborne fractions of *Chenopodium* collection was investigated and assessed. The outcome demonstrated that the $(\text{CH}_3)_2\text{CO}$ concentrate of *Chenopodium* collection (ACCA) has indicated huge decrease in rodent paw edema (80.13%) at measurements level of 200mg/kg orally in 21 days of the investigation. On 22nd day, it was watched that the modified the parameters of blood (i.e. Hb, WBC, RBC & ESR), biochemically active markers (serum creatinine, add up to proteins & intense stage proteins) and misfortune in weight of the body in ligament rats were

fundamentally taken back to close ordinary level by the ACCA separate. ACCA remove fundamentally diminished the NFkB articulation in paraventricular core of hypothalamus and this impact is practically identical with standard indomethacin. [20]

3.4. Analgesic effect

Significant analgesic activity was revealed by crude extract (at dose of 500 mg/kg from 30 min - 210 min against tail flick method in mice. [21]

3.5. Spasmolytic effect

The plant extraction in ethanol and fractionated in ethyl acetic acid derivation, chloroform, n-butanol & water was done. The rough concentrate and its portions were tried in vitro on smooth muscles of intestine in rabbit. The rough concentrate showed a dosage subordinate increment in unwinding of smooth muscles, beginning with 5 mg/ml and most extreme impact was initiated at 20 mg/ml (92.86%). Every one of the portions were added to rabbit's digestive tract at 15 mg/ml dosage. The ethyl acetic acid derivation & chloroform parts of *Chenopodium* collection displayed unwinding of the muscles of the intestine (43.48 and 51.52%, separately); while, n-butanol division of *Chenopodium* collection delivered solid relaxant impact (91.18%). [22]

3.6. Gastroprotective effect

The impact of alcoholic concentrate of *Chenopodium album* was researched using rats to assess the ulcer healing action by utilizing 3 animal models including "pyloric ligation, ethanol and chilly limitation push-actuated ulcers". Alcoholic concentrate essentially diminishes the volume of gastric corrosive emission, free causticity; add up to sharpness and ulcer record as for control. Areas of ulcerated territory uncovered to be increased in the width of recovered glandular-epithelium after treatment with the liquor extricate. The collagen's content within ulcerated tissue was fundamentally expanded by liquor concentrate and ranitidine as positive control. No noteworthy distinction on slim thickness in scar tissue was seen after treatment with liquor concentrate or ranitidine. [23]

3.7. Hepatoprotective effect

The cell reinforcement and liver healing viability of the *chenopodium album* separate (300 mg/kg & 450 mg/kg) was assessed in carbon tetra-chloride (CCl_4) incited toxicity of liver in rats. *Chenopodium album* remove was found to display astounding cancer prevention agent and free radical searching movement, when contrasted and ascorbic corrosive, in vitro considers, *chenopodium album* separate at a dosage of 450 mg/kg demonstrated restraint of lifted biochemical markers related with enlistment of hepatotoxicity by CCl_4 . These were additionally lessened histopathologic impacts of CCl_4 . [24]

In another study, the alcoholic & water extracts of all aerial parts of *Chenopodium album* (at the doses of 200 & 400

mg/Kg) were tested for liver protective potential using paracetamol induced liver toxicity model. The aqueous extract at a dosage of 400 mg/kg was observed to be stronger when contrasted with Silymarin. The alcoholic and watery concentrates of *Chenopodium* collection essentially reestablish physiological uprightness of hepatocytes. Fluid and alcoholic concentrate did not hint at any harmfulness up to oral dosage of 5 g/Kg in mice. [25]

Literature also revealed the hepatoprotective exercises of dried entire plant of *Chenopodium* collection Linn, ethanol and methanol removes in proportion of (50:50), was likewise assessed against paracetamol prompted hepatic damage. (CH₃)₂CO and methanol separate at a measurement of 400mg/kg orally, demonstrated huge ($p < 0.001$) liver protective action, their impact was like the standard medication, silymarin. [26]

3.8. Anticancer effect

Treatment with extracts in different solvent i.e. ethyl acetate, petroleum ether & methanolic leaves extract of *Chenopodium album* was observed on the growth of estrogen dependent (i.e. MCF-7) & estrogen-independent (i.e. MDA-MB-468) human breast cancer cell lines using trypan blue exclusion and MTT [3-(4, 5-dimethyl thiazol-2-yl)-2, 5-diphenyl tetrazolium] bioassay technique. Methanolic leaves extract of *Chenopodium album* unveiled extreme anti-breast cancer efficacy with IC₅₀ value 26.91 mg/ml against MCF-7 cell line. Significant percent inhibition of 94.06% was noted for methanolic leaves extract of *Chenopodium album* at 48th hour of acquaintance & the concentration 100 mg/ml ($p < 0.05$) against MCF-7 "breast cancer cell line". [27]

3.9. Effect on male reproduction

Ethanollic concentrate of *Chenopodium* collection at measurements of 100, 250 and 500mg/kg orally, in male pale skinned person mice demonstrated huge increment in the mount recurrence, intromission recurrence, intromission idleness and total of penile reflexes and huge diminishment in the post ejaculatory interim. Also 500 mg/kg, orally, was observed to be the best measurement. The ethanollic concentrate of seeds of *Chenopodium* collection was assessed for its impact on anabolic action, sexual conduct and sperm check in male rats. Organization of ethanollic extricate at a grouping of 200 mg/kg bw brought about articulated anabolic impact in regarded creatures as prove by an expanded body weight and additionally the heaviness of conceptive organs. Sexual conduct and execution were likewise especially enhanced as reflected in decrease of mount, intromission and post ejaculatory idleness. Besides, the concentrate additionally improved sperm check. Nonetheless, then again, the impact of *Chenopodium* collection seed extricate (CAE) instigated sperm demise, the impact which is expected to (a) lipid peroxidation of the sperm cell layer, oxidation of some basic cell proteins and consumption of intracellular diminished

gluthathione, demonstrating creation of ROS; (b) actuation of Mn-SOD and inactivation of catalase favoring endogenous gathering of H₂O₂; (c) age of O₂-at an improved rate amid oxidative worry as confirm by expanded Mn-SOD action and protein articulation; (d) aggregation of ROS in spermatozoa and (e) expanded generation of O₂-and H₂O₂ prompted apoptosis-like passing in sperm cells as saw by DNA stepping stool development. In this way, the sperm demise caused by CAE is because of oxidative harm of cell macromolecules by in situ age of ROS. Aqueous decoction of *Chenopodium album* seeds was surveyed for its "sperm-immobilizing" and preventative adequacy in research center warm blooded animals. The base successful centralization of CAD that incited prompt immobilization of rodent spermatozoa in vitro was 2 mg/ml. The component of CAD activity included breaking down of sperm plasma layer and disintegration of acrosomal top causing sperm demise. Preparation of oocytes and foundation of embedding were counteracted in the uterine portion that was directed with seed extract. In rabbit, intravaginal utilization of extract altogether obstructed the foundation of pregnancy. In like manner, CAD has obvious spermicidal potential, which might be investigated as an effective component of vagina prophylactically. [28]

3.10. Contraindication and adverse effects

Chenopodium album was an allergenic plant. A few arrangements of its concentrates were utilized for determination and immunotherapy of patients. The hypersensitive concentrate of *Chenopodium* collection dust has been arranged and inspected in skin prick testing in correlation with a business item in Iran. [29]

The impacts of a concentrate arranged from *Chenopodium album* dust were explored to actuate hypersensitive asthma in BALB/C mice. Mice were sharpened by ip infusion and an intratracheal instillation of the concentrate of *Chenopodium* collection. *Chenopodium* collection remove expanded serum levels of particular IgE and creation of IL-4 and IL-5 from splenocytes. An aviation route eosinophilia was likewise exhibited in mice. Sun presentation after oral admission of *Chenopodium* collection can prompt sunburn-like rashes inferable from its furocoumarin content. Numerous investigations recorded that patients created dermatitis with edema, erythema and rot on the face and dorsum of the hands when they presented to daylight in the wake of eating *Chenopodium album*.

The safety standard tool of "*Chenopodium album* seed decoction (CAD)" was assessed. *In vitro* bothering ponders in rabbit's erythrocytes uncovered the blood diminishing record of "*Chenopodium album* seed decoction (CAD)" to be 8.2 mg/ml. The dermal aggravation test demonstrated that the plant wasn't aggravation even at higher dosages. Intra vaginal utilization of "*Chenopodium album* seed decoction (CAD)" in rodent vagina for 14 sequential days caused slight reversible aggravation on vaginal epithelial cells at measurements as high

as 82 mg/ml. Be that as it may, at this measurement level it neither had any antagonistic impact on vaginal tissue expansion nor did it cause in situ apoptosis as obvious from PCNA recoloring and TUNEL test. Ripeness and fertility were reestablished 4-15 days after withdrawal of “Chenopodium album seed decoction (CAD)” application.

Nevertheless, *Chenopodium album* may result in lethal intoxication in ruminants because it accrued high nitrate levels as well as it may also be accumulated as the soluble oxalate. The nitrate-nitrogen levels of 2,500 ppm were assumed to be associated with mortality in cattle in *Chenopodium album* plant. [29]

4. CONCLUSION

The present study concluded as *Chenopodium album* to be most promising plant having distinguished medicinal values with extensive variety of pharmacological potentials that could be consumed in quite a lot of remedial purposes owing to its usefulness and safety.

Chenopodium album is great wellspring of practical supplements and has restorative properties. It can be joined in various expelled sustenance items to make them more nutritious, more advantageous and in addition shopper arranged. The expansion of its leaves to expelled items can upgrade the synthetic and wholesome parameters and can enhance expelled items as practical nourishments. The plants likewise have high natural exercises thus might be of awesome therapeutic esteem. Business abuse of *Chenopodium album* in numerous districts of the world is still a long way from reality. The dynamic constituents can be disengaged and additionally assessed for the advancement of helpful medications. Their cancer prevention agent and antibacterial exercises additionally loan trustworthiness to the natural estimation of this plant. These trials should clear route for the utilization of *Chenopodium album* in districts where the green verdant vegetables are developed however yet to perceive any business abuse. Expanded mindfulness in the general public and subsequently more utilization of this plant may go long path towards forestalling not just lack ailments and age related strong degeneration related issue, yet in addition secure against incessant degenerative illnesses, for example, malignancy and cardiovascular issue which eventually will be exceptionally beneficial to the rural community.

5. REFERENCES

- Chenopodium album, http://www.nps.gov/akso/natres/epmt/Species_bios/Chenopodium%20album.pdf.
- Agrawal MY, Agrawal YP and Shamkuwar PB. *Int J PharmTech Res*, 2014; **6(1)**:383-391.
- Wiat C. Medicinal plants of the east pacific. Drugs for the Future? World Scientific Publishing Co Pte Ltd, 2006, 100.
- Pande M and Pathak A. *Asian J Exp Biol Sci*, 2010; **1(1)**: 91-95.
- Singh PK, Dwevedi AK and Dhakre G. *International Journal of Applied Biology and Pharmaceutical Technology*, 2011; **2(3)**: 398-401.
- The earth of India. All about Bathua (*Chenopodium album*) 2014, <http://theindianvegan.blogspot.com/2014/01/all-about-bathua-chenopodium-album.html>
- Holm, Leroy G, Plucknett DL, Pancho JV and Herberger JP. The world's worst weeds: distribution and biology. East-West Center, University Press of Hawaii, 1977, 84-91.
- Arora SK, Itankar PR, Verma PR, Bharne AP and Kokare DM. *J Ethnopharmacol*, 2014; **155(1)**: 222-229.
- Agarwal SS, Yamrekar BP and Paridhavi M. Clinical useful herbal drug. Ahuja Publishing House, New Delhi, 2005, 10-12.
- Panda H. Handbook on medicinal herbs with uses. Asia Pacific Business Press, New Delhi, 2005, 325-326.
- Pramila K, Neetu S and Anju R. *Indian J Traditional Knowledge*, 2006; **5(3)**: 300-309.
- Khare CP. Indian medicinal plants. Springer International Publication, New Delhi, 2007, 141-142.
- Priya S, Yogesh S, Singhai AK and Abhishek S. *Research Journal of Pharmacy and Technology*. 2010; **3(4)**: 960-963.
- Baldi A and Choudhary NK. *IJGP*; **7(1)**: 50-56.
- Gogoi B and Zaman K. *Journal of Pharmacognosy and Phytochemistry*, 2013; **2(2)**:30-40.
- Pal A, Banerjee B, Banerjee T, Masih M and Pal K. *International Journal of Pharmacy and Pharmaceutical Sciences*, 2013; **3**: 55-57.
- Yogesh S, Priya S, Utkarsh U, Sumit S, Shivakant S, Singhai AK, and Prashant S. *Pharmacognosy Journal*, 2010; **2(14)**:7-10.
- Adedapo A, Jimoh F and Afolayan A. *Acta Pol Pharm*, 2011; **68(1)**: 83-92.
- Sood P, Modgil R, Sood M and Chuhan PK. *Annals Food Science and Technology*, 2012; **13(1)**: 68-74.
- Rastogi RP and Mehrotra BN. Compendium of Indian medicinal plants. Vol. 3. Reprint ed. CDRI, Lucknow, 1998, 162-163.
- Cutillo F, D'Abrosca B, Della Greca M and Zarrelli A. *Chem Biodivers*, 2004; **1(10)**: 1579-1583.
- Cutillo F, Della Greca M, Gionti M, Previtera H and Zarrelli A. *Phytochemical Anal*, 2006; **17(5)**: 344-349.
- Jhade D, Padmaa MP and Usha G. *J of Pharmacy Res*, 2009; **2(7)**: 1192-1193.
- Della Greca M, Di Marino C, Zarrelli A and D'Abrosca B. *J Nat Prod*, 2004; **67(9)**: 1492-1495.
- Lavaud C, Voutquenne L, Bal P and Pouny I. *Fitoterapia*, 2000; **71(3)**: 338-340.
- Nahar L and Sarker SD. *Brazilian Journal of Pharmacognosy*, 2005; **15(4)**: 279-282.
- Dutt S, Yadav OP, Kapoor HC and Lodha ML. *Indian J Biochem Biophys*, 2004; **41(1)**: 29-33.
- Barderas R, Villalba M, Lombardero M and Rodríguez R. *Int Arch Allergy Immunol*, 2002; **127(1)**: 47-54.
- Dai Y, Ye WC, Wang ZT, Matsuda H, Kubo M and But PP. *J Ethnopharmacol*, 2002; **81(2)**: 2002, 245-250.