



A Review on *Digera Muricata* (L.) Mart. - A Great Versatile Medicinal Plant

Neha Sharma*, Rekha Vijayvergia

Plant Pathology and Plant Biochemistry Laboratory, Department of Botany, University of Rajasthan, Jaipur, India.

*Corresponding author's E-mail: siya28@gmail.com

Accepted on: 05-03-2013; Finalized on: 30-04-2013.

ABSTRACT

Digera muricata (L.) Mart. is an annual herb, growing to 20-70 cm tall. It is an important medicinal herb belongs to the Amaranthaceae family, found as a weed throughout India. Though almost all of its parts are used in traditional systems of medicines, leaves, roots and shoots are the most important parts which are used medicinally. The present article gives an account of updated information on its phytochemical and pharmacological properties. The review reveals that wide numbers of phytochemical constituents have been isolated from the plant which possesses activities like antibacterial, antifungal, diuretic, laxative, Free radical scavenging activity, anthelmintic, and various other important medicinal properties. The crushed plant is used as mild astringent in bowel complaints. The Leaves and spikes are used as a vegetable. Flowers and seeds used in the treatment of urinary discharges. For the last few decades or so, extensive research work has been done to prove its biological activities and pharmacology of its extracts. Analysis of various fractions of the *D. muricata* indicated the presence of flavonoids, alkaloids, terpenoids, saponins, coumarins, tannins, cardiac glycosides and anthraquinones.

Keywords: *Digera muricata*, Amaranthaceae, pharmacological activities, medicinal properties, phytochemical.

INTRODUCTION

From the time immemorial, plants have been widely used as curative agents for variety of ailments. Any plant which possesses curative elements or properties in one or more of its organs may be termed as medicinal plant. Plants based medicaments have been employed since the dawn of civilization for prolonging the life of man and for combating various ailments.

Medicinal herbs are moving from fringe to mainstream use with a greater number of people seeking remedies and health approaches free from side effects caused by synthetic chemicals. India officially recognizes over 3000 plants for their medicinal value. It is generally estimated that over 6000 plants in India are in use in traditional, folk and herbal medicine.

Indian traditional medicine is based on various systems including Ayurveda, Siddha, Unani and Homoeopathy. The evaluation of these drugs is primarily based on phytochemical, pharmacological and allied approaches including various instrumental techniques such as chromatography, microscopy and others. With the emerging worldwide interest in adopting and studying traditional systems and exploiting their potential based on different health care systems, the evaluation of the rich heritage of traditional medicine is essential². In this regard, one such plant is *Digera muricata*.

Digera is a genus having one species only *Digera muricata* (L.) Mart. (Syn *Digera arvensis* Forssk ; *Achyranthes muricata* L.) belongs to the family amaranthaceae. This is an annual herb, growing to 20-70cm tall. It is widespread in eastern tropical Africa and subtropical Asia. In India, It is widely distributed in Rajasthan, Maharashtra and Andhrapradesh. The root,

leaf, stem, seeds and flowers of this plant have medicinal properties and traditionally used as medicinal plant. All parts of the plant have been used as crude drug for the treatment of kidney stone and urinary tract disorders.

D. muricata ethnopharmacologically has been used in renal disorders, aperients, refrigerant³. This plant is also used as an alternative for secondary infertility⁴. Antioxidant properties of *D. muricata* against the CCl₄-induced toxicity for kidneys and testis had been documented. The Leaves and young shoots of this plant are locally used as a vegetable and given to relieve constipation⁵. *D. muricata* is used internally against digestive system disorders and in India seeds and flowers are used to treat urinary disorders. Leaf paste is applied locally to prevent pus formation⁶. This article aims to provide a comprehensive review on the phytochemical and pharmacological aspects of *Digera muricata* (L.) Mart.

TAXONOMY

Kingdom	: Plantae (plants)
Subkingdom	: Tracheobionta (Vascular plants)
Superdivision	: Spermatophyta (Seed plants)
Division	: Magnoliophyta (flowering plants)
Class	: Magnoliopsida (Dicotyledons)
Subclass	: Caryophyllidae
Family	: Amaranthaceae (Amaranth)
Order	: Caryophyllales
Genus	: <i>Digera</i> Forssk
Species	: <i>Muricata</i> (False amaranth)
Subspecies	: <i>Digera muricata muricata</i>
Subspecies	: <i>Digera muricata trinervis</i>
Variety	: <i>Digera muricata macroptera</i>



Variety : *Digera muricata muricata*
 Variety : *Digera muricata patentipilosa*

Botanical Name

Digera muricata (L.) Mart.

Synonyms

Achyranthes alternifolia L., *Achyranthes muricata* L.,
Digera alternifolia (L.) Aschers., *Digera arvensis* Forssk.

Common name:

Hindi : Latmahuria, Lesua
 Sanskrit : Aranya, Aranyavastuka, kunanjara, kuranjara
 English : False amaranth
 Telugu : Chnchali Koora
 Tamil : Toya Keeri, kaatu Keerai
 Kannada: Chenchali soppu, Goraji playa, Kankali soppu
 Marathi : Gitana, Getna
 Bangali : Lata mouri Ful, Gun gutiya

Distributional Range

This weed flower is known as false amaranth. It is widespread in eastern tropical Africa (from Sudan and Ethiopia south to Tanzania)⁷, Madagascar⁸ and tropical and subtropical Asia (from Yemen to Afghanistan, Pakistan, India, Malaysia and Indonesia). In India it is widely distributed in Rajasthan, Maharashtra and Andhrapradesh.

Native

- **Africa**
 Northeast Tropical Africa: Ethiopia; Somalia; Yemen - Socotra
 East Tropical Africa: Kenya; Tanzania; Uganda
- Western Indian Ocean: Madagascar
- **Asia-temperate**
 Arabian Peninsula: Oman; Saudi Arabia; Yemen
 Western Asia: Afghanistan; Iran
- **Asia-tropical**
 Indian Subcontinent: India; Pakistan
 Malaysia: Indonesia -Celebes, Java, oluccas; Malaysia

Cultivated:

- **AFRICA**
 Northeast Tropical Africa: Ethiopia
- **ASIA-TROPICAL**
 Indian Subcontinent: India

Ecology

D. muricata is most common on disturbed and waste ground, but occurs in many kinds of habitat, from dry savanna and semi-desert to moist localities on deep clay and mud soils, from sea-level up to 1500 m altitude. It also occurs as a weed in fields, sometimes being

troublesome. Its cultivation occurs in northeast tropical Africa (Ethiopia) and in Indian subcontinent (India)⁹.

Nutritive Value

D. muricata is considered as an edible GLV (Green leafy vegetable)¹⁰. Fifty six percent edible portions are present in this weed¹¹. This plant is a rich source of calcium, iron, phosphorus, potassium, magnesium etc. Various parameters are listed in table.

Table 1: Nutrient levels of *Digera muricata*

Parameter	Concentration (g/100g)
Edible portion	56
Ash value	3.54
Moisture	83.8
Protein	4.3
Mineral contents	mg/100g
Calcium	506
Potassium	604
Magnesium	232
Sodium	-
Phosphorus	63
Trace mineral contents (mg/100g)	
Iron	17.72
Zinc	0.57
Copper	0.16
Chromium	0.243
Manganese	0.23
Vitamin content (mg/100g)	
Ascorbic acid	49
Thimine	0.10
Total-Carotene	17.93
B-carotene	3.36

BOTANICAL DESCRIPTION

This is an annual herb up to 70 cm tall; stem simple or branched, subglabrous, ridged. Leaves alternate, simple; petiole up to 5 cm long; blade linear to ovate, 1–9 cm × 0.2–5 cm, base narrowed, apex acuminate, margin entire, subglabrous. Inflorescence a long-pedunculate (up to 14 cm long), axillary, spike-like bracteate raceme up to 30 cm long, each bract subtending a subsessile partial inflorescence with a central fertile flower and 2 sterile lateral flowers. Flowers are borne on slender spike-like racemes, which can be as large as 30 cm long. Flowers are hairless, white mixed with pink to carmine or red, usually becoming greenish-white in fruit. Flowering occurs in month of August and September. Fertile flower with 2 firm, boat-shaped outer perianth segments 3–5 mm long and 2–3 inner, slightly shorter, hyaline segments; stamens usually 5, free or slightly connate at base; ovary superior, 1-celled, style filiform, up to 4 mm long, stigmas 2, divergent; lateral flowers consisting of accrescent antler-shaped scales. Fruit is subglobose, hard, 2 mm in

diameter, ridged, enclosed by the persistent perianth and falling together with the sterile flowers and bracteoles.

Digera comprises only 1 species. Based on the venation of the outer tepals 2 subspecies of *Digera muricata* have been distinguished: subsp. *muricata* with outer tepals 7–12-veined, mainly occurring in Asia, but also in eastern Africa and Madagascar, and subsp. *trinervis* C.C.Towns. with outer tepals 3–5-veined, mainly occurring in Africa. Based on hairiness of leaves and on form of scales in sterile flowers, several varieties have been distinguished in subsp. *trinervis*, of which var. *patentipilosa* C.C.Towns. It seems most suitable as a leafy vegetable because it has large leaves¹².



Figure 1: a. A complete plant of *Digera muricata* (L.) Mart. b. Leaves c. Inflorescence

MEDICINAL PROPERTIES

The *Digera muricata* (L.) Mart. is a wild edible herb used by village people. It is popularly known for herbal remedy for various ailments. In Ayurveda this herb is considered as cooling, astringent of bowels and also used as a laxative¹³. The leaves are used for treatment of diabetic¹⁴. But the scientific basis for its medicinal use especially for boiled root infusion given to mother after child birth to increase lactation purpose is to be evaluated. The flower and seeds are used to treat urinary discharges¹⁵. Ethyl alcohol extract of plant is diuretic. The whole plant is used in digestive system disorders. The leaves and young shoots of this plant are locally used as a vegetable and given to relieve constipation¹¹. The whole plant is used in urinary disorders¹⁶. The decoction of leaves gives once in a day for kidney stone treatment^{17,18}. The extract of this plant used in biliousness and in urinary discharges¹⁹.

Leaf paste is applied locally to prevent pus formation. The crushed plant is used as mild astringent in bowel complaints and antibilious²⁰. Antioxidant properties of *D. muricata* against the CCl₄-induced toxicity for kidneys and testis had been documented²¹. This plant is used as an alternative for secondary infertility. Secondary infertility is found to be associated with hepatic disorders. The models created by the use of CCl₄ to induce liver injuries can be best suited to study the hypogonadism in rat. The whole plant extract improves blood content and also works as expectorant²². This is antiperiodic, coolent and stomachin²³.

Other uses

D. muricata is considered as a famine food²⁴ because of rich source of nutrients²⁵. In Kenya they are particularly popular as a cooked vegetable amongst coastal tribes. In India the leaves are made into curries or the entire plant is boiled in water and seasoned with salt and chilli. The whole plant is also commonly grazed as forage, particularly by sheep and goats. The flowers are rich in nectar which is sometimes sucked by children in Kenya²⁶.

CHEMICAL CONSTITUENTS

The primary metabolites like carbohydrates, proteins, lipids, phenols, chlorophylls, amino acids etc. of this plant in different solvent extracts have been investigated^{27,28}. The plant contains α - and β - spinasterol²⁹. Analysis of various fractions of the *D. muricata* indicated the presence of flavonoids, alkaloids, terpenoids, saponins, coumarins, tannins, cardiac glycosides and anthraquinones. Rutin and Hyperoside flavonoids have been identified in hexane extract of this plant³⁰.

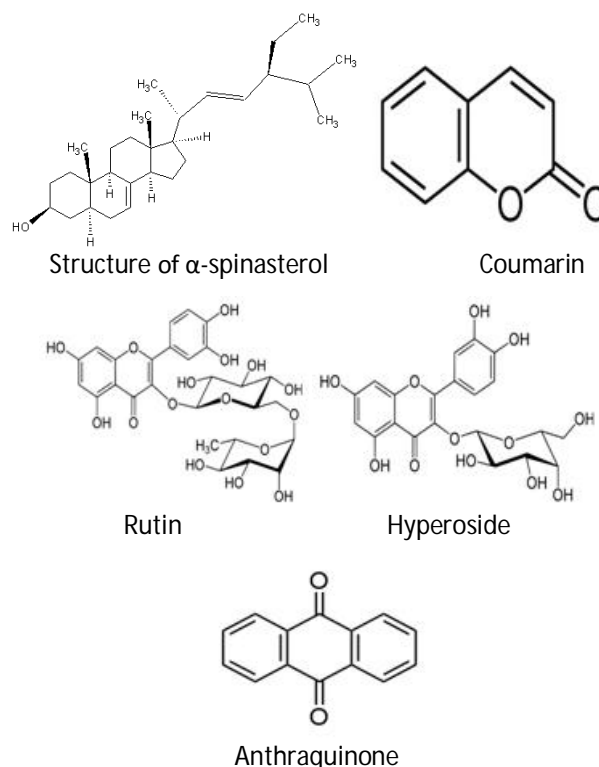


Figure 2: Structure of some phytoconstituents isolated from *Digera muricata*

PHARMACOLOGICAL ACTIVITIES

Hepatoprotective activity

The methanolic extract of *D. muricata* shows hepatoprotective effect against acryl amide-induced hepatocellular injuries. Acryl amide (AA) is a water-soluble vinyl monomer used in the production and synthesis of polyacrylamides^{31,32}. It has been documented that AA is formed during the cooking of starchy foods at high temperature³³. Daily exposure to AA might present a risk factor for neurotoxicity and reproductive toxicity as well as carcinogenicity in humans^{34,35}. AA can also cause glutathione depletion, resulting in intracellular oxidative stress³⁶. The methanolic extract of *D. muricata* was given to acryl amide induced Sprague-Dawley rats and found that Hepatic lesions induced with AA were reduced with DME treatment. The results suggest that the hepatoprotective effects of DME against AA-induced oxidative injuries could be attributed to the phenolics and flavonoids.

Antimicrobial activity

The different solvent extracts shows antifungal and antibacterial activity against selected bacteria and fungi. The organic successive Soxhlet extracts of *D. muricata* i.e., petroleum ether, chloroform, ethanol and distilled water, have shown significant zone of inhibition of bacterial growth at the concentrations of 200 and 400 µg/well against test pathogens²⁸. It is also reported that the methanol extract shows maximum activity against test bacteria and fungi²⁷.

Antioxidant activity

The plant has shown antioxidant activity in different investigations. Mety et al., 2011 analysed free radical scavenging and antioxidant activity of different solvent extract like hexane, petroleum ether, chloroform, methanol, ethanol and aqueous³⁷. The maximum activity recorded in methanol and least activity was recorded in hexane. The methanolic crude extracts of *D. muricata* was screened for their free radical scavenging properties by DPPH (1,1-diphenyl-2-picryl hydrazyl) radical scavenging assay. The maximum activity was observed in roots of *D. muricata*³⁸. Antioxidant properties of *Digera muricata* methanol extract against the CCl₄-induced toxicity in kidneys and testis had been well documented^{39,21}.

Anti-diabetic activity

The methanolic extract of *D. muricata* (MEDM) leaves exhibited antidiabetic activity in alloxan induced diabetic rats. These results suggest that MEDM (200mg/kg) showed antihyperglycemic activity in diabetic rats. The other parameters like blood glucose level, HDL level in serum decreases and body weight increases¹⁴.

Anthelmintic activity

The crude extract from leaves was preliminary screened for anthelmintic activity when tested against earthworms (*Pheretima posthuma*)⁴⁰.

Anti-testicular toxicity

The study suggested the protective potential of hexane extract of *D. muricata* against the CCl₄-induced liver and testicular toxicity. CCl₄ can rapidly lead to both oxidative stress and acute liver injuries^{41,42}. Liver cirrhosis causes Hypogonadism in male rats which are cured by Hexane extract of *D. muricata*. DMH treatment ameliorated the hepatic injuries with consequent increase in the antioxidant status of various enzymes and compounds. Level of testosterone was elevated with DMH in addition to the repairing of testis and accessory organs. These protective effects of DMH against the CCl₄ toxicity may be attributed due to the presence of various bioactive groups and specifically the rutin and hyperoside in DMH⁴³.

Renal disorders

D. muricata is used in renal disorders in folk medicine. The extract of this plant is administered daily in kidney stone treatment⁴⁴. Generation of reactive radicals has been implicated in carbon tetrachloride-induced nephrotoxicity, which are involved in lipid peroxidation, accumulation of dysfunctional proteins, leading to injuries in kidneys. Nephrotoxicity is a poisonous effect of some substances on kidneys. The n-hexane and methanolic extract of *D. muricata* shows protective role against Carbon tetrachloride which is induced nephrotoxicity in rats³⁹.

Allelopathic effect

The aqueous extract of stem, root and leaf of *D. muricata* shows allelopathic effect on in vitro seed germination of *Pennisetum typhoideum* (bajra). Different concentrations of various parts of weed showed inhibitory effects on shoot and root growth of *Pennisetum typhoideum*. The leaf extract proved inhibitory in nature than stem and root⁴⁴.

Protective effect

The methanolic and hexane extract of *D. muricata* shows protective effect against oxidative stress caused by ccl4 in rats⁴⁵. It is able to ameliorate oxidative stress in adrenal gland induced by CCl₄ in rat. The protective potential may also involve the preventive effects of *D. muricata* methanolic extract by the inhibition of CCl₄ metabolism⁴⁶. This study further supports the scientific evidence in favor of its pharmacological use in oxidative stress diseases.

DISCUSSION

Before the introduction of modern medicines, disease treatment was entirely managed by herbal remedies. It is estimated that about 80% of the world population residing in the vast rural areas of the developing and under developed countries still rely mainly on medicinal



plants. Phytochemical and pharmacological investigations carried out in the plant reveals its multidisciplinary usage. It is quite obvious that the plant is widely used in traditional medicinal system of India and has been reported to possess anti-bacterial, antifungal, anti-diabetic, hepatoprotective, nephrotoxicity protective, anthelmintic, free radical scavenging properties. It is known as a rich source of phenols, tannins, terpenoids, flavonoids and glycosides present in *Digera muricata* might be medicinally important and/or nutritionally valuable. The plant is rich in carbohydrates, Calcium, potassium, ascorbic acid, iron, magnesium etc. The present review summarizes some important pharmacological studies on *D. muricata* and phytochemical investigations and isolated principles from them, which can be investigated further to achieve lead molecules in the search of novel herbal drugs.

REFERENCES

1. Padulosi, DD, Leaman and F. D. Quick 2002. Challenges and opportunities in enhancing the biodiversity conservation and uses of medicinal plants. Herbs, spices and medicinal plants, 9, 243-279.
2. Gupta RK, Medicinal & Aromatic plants, CBS publishers & distributors, 1st edition, 2010, 116-117.
3. Anjaria J, Parabia M, Bhatt G and Khamar RA, glossary of selected indigenous medicinal plants of India. SRISTI Innovations PO Box: 15050, Ahmedabad-380 015, India, 2002, 26.
4. Hocking GM. Pakistan medicinal plants-IV. *Qualitas Plantarum* 9, 2, 1962, 103-119.
5. Chettleborough J, Lumeta J and Magesea S, Community use of nontimber forest products: A case study for the Kilombero valley. Integrated Environmental Programme. The Society for Environmental Exploration UK and the University of Da es Salaam. Frontier Tanzania, 2000, 16.
6. Katewa SS, Chaudhary BL and Jain A, Folk herbal medicines from tribal area of Rajasthan, India, *Journal of Ethnopharmacology*, 92, 2004, 41-46.
7. Townsend, CC, Amaranthaceae. In: Edwards, S, Mesfin Tadesse, Demissew Sebsebe & Hedberg, I. (Editors), *Flora of Ethiopia and Eritrea. Volume 2, part 1. Magnoliaceae to Flacourtiaceae*. The National Herbarium, Addis Ababa University, Addis Ababa, Ethiopia and Department of Systematic Botany, Uppsala University, Uppsala, Sweden, 2000, 299-335.
8. Cavaco A, Amaranthacées (Amaranthaceae). *Flore de Madagascar et des Comores (plantes vasculaires)*, familles 66-69. Firmin-Didot et cie., Paris, France, 1954, 56 pp.
9. Schippers RR, African indigenous vegetables. An overview of the cultivated species. Natural Resources Institute/ACP-EU Technical Centre for Agricultural and Rural Cooperation, Chatham, United Kingdom, 2000.,214.
10. Seshadri S & Nambiar, VS, Kanjero (*Digera arvensis*) and Drumstick leaves (*Moringa oleifera*): nutrient profile and potential for human consumption. In: Simopoulos, A.P. & Gopalan, C. (Editors). *Plants in human health and nutrition policy*. *World Review of Nutrition and Dietetics*, 91, 2003, 41-59.
11. Gupta SA, Jyothi Lakshmia, Manjunathb MN, Jamuna Prakasha, Analysis of nutrient and antinutrient content of underutilized green leafy vegetables, *LWT*, 38, 2005, 339-345.
12. Townsend CC, Amaranthaceae. In: Polhill, R.M. (Editor). *Flora of Tropical East Africa*. A.A. Balkema, Rotterdam, Netherlands, 1985, 136.
13. Parrota JA, Haeling plants of Pennninsular india, CABI publishing, CAB international Newyork, USA, 2001, 56.
14. Jagatha G and Senthilkumar N, Evaluation of anti-diabetic activity of methanol extract of *Digera muricata* (l) Mart in alloxan Induced diabetic rats, *International journal of pharmaceutical science and research*, 2(6), 2011, 1525-1529.
15. Rajasab AH and Isaq M, Documentation of folk knowledge on edible wild plants of north Karnataka. *International journal of traditional knowledge*. 3(4), 2004, 419-429.
16. Qureshi R and Bhatti GR, folklore uses of Amaranthaceae family from Nara desert, Pakistan, *Pak. J. Bot.*, 41(4), 2009, 1565-1572.
17. Aggarwal S, Gupta V and Narayan R, Ecological studies of wild animal plants in a dry tropical peri-urban region of Utter Pradesh in India, *Int. J. Med. Arom. Plants.*, 2(2), 2012, 246-253.
18. Sharma N, Tanwer BS and Vijayvergia R, Study of medicinal plants in Aravali regions of Rajasthan for treatment of Kidney stone and Urinary tract troubles. *International Journal of PharmTech Research*, 3(1), 2011, 110-113.
19. Khare CP, An Indial medicinal plants. An illustrative dictionary, Springer, 2008, 213.
20. Rahman MA, Khudeja B, Rashid ME and Rashid MH, Medicinal plant diversity in the flora of Bangladesh and their conservation: 2. a report on ten angiosperm families. *Plant Archives*. 12(2), 2012, 1023-1035.
21. Khan MR and Ahmed D, Protective effects of *Digera muricata* (L.) Mart. on testis against oxidative stress of carbon tetrachloride in rat. *Food Chem. Toxicol.*, 47, 2009, 1393-1399.
22. Shah A, Marwat SK, Gohar F, Khan A, Bhatti KH, Muhammad A4, Din NUD, Ahmad M, Zafar M, Ethnobotanical Study of Medicinal Plants of Semi-Tribal Area of Makerwal & Gulla Khel (Lying between Khyber Pakhtunkhwa and Punjab Provinces), Pakistan, *American Journal of Plant Sciences*, 4, 2013, 98-116.
23. Kumar ACK, Revathi K, Mohanalakshmi S, A review on edible herbs as haematinics. *International Journal of Pharmacy*, 2(2), 2012, 44-53.
24. Freedman, R.L., 1998. *Famine foods: Amaranthaceae*. [Internet] Purdue University, West Lafayette, Indiana, United States. [www.hort.purdue.edu/newcrop/ Famine Foods/ ff_families/AMARANTHACEAE.html](http://www.hort.purdue.edu/newcrop/Famine_Foods/ff_families/AMARANTHACEAE.html). Accessed August 2003.
25. Benson AM, Hunkeler MJ, Talalay P, Increase of NADPH, quinone reductase activity by dietary antioxidant: Possible role in protection against carcinogenesis and toxicity. *Proceedings of the National Academy of Sciences of the United State of America*, 77, 1980, 5216-5220.

26. Maundu, PM, Ngugi, GW & Kabuye, CHS, Traditional food plants of Kenya. Kenya Resource Centre for Indigenous Knowledge (KENRIK), Nairobi, Kenya, 1999, 270.
27. Sharma N, Tanwer BS and Vijayvergia R, Study of primary metabolites and antimicrobial activities of *Digera muricata* (L.) Mart., J. Chem. Pharm. Res., 2011, 3(2), 424-431.
28. Mathad P and Mety SS, Phytochemical and Antimicrobial Activity of *Digera muricata* (L.) Mart. E-Journal of Chemistry, 2010, 7(1), 275-280.
29. Pullaiah T. The Encyclopedia of World Medicinal Plants. Regency publishers. 3, 2006,776.
30. Khan MR, Khan GN and Ahmed D, 2011, Evaluation of antioxidant and fertility effects of *Digera muricata* in male rats, Asian journal of pharmacy and pharmacology, 5(6), 688-699.
31. Paulsson B, Granath F, Grawe J, Ehrenberg L, Tornqvist M, The multiplicative model for cancer risk assessment: applicability to acryl amide. Carcinogenesis, 22, 2001, 817-819.
32. Friedman M, Chemistry, biochemistry, and safety of acrylamide. A review. J. Agri. Food Chem., 51, 2003, 4504-4526.
33. Taubert D, Harlfinger S, Henkes L, Berkels R, Schomig E, Influence of processing parameters on acrylamide formation during frying of potatoes, J. Agric. Food Chem. 52, 2004, 2735-2739.
34. Svensson K, Abramsson L, Becker W, Glynn A, Hellena s KE, Lind Y, Rose'n J, Dietary intake of acrylamide in Sweden. FoodChem. Toxicol. 41, 2003, 1581-1586.
35. Klaunig JE. Acrylamide carcinogenicity. J. Agric. Food Chem. 56, 2008, 5984-5988.
36. Tong GC, Cornwell WK, Means GE, Reactions of acrylamide with glutathione and serum albumin. Toxicol. Lett., 147, 2004, 127-131.
37. Mety SS, Mathad P and Rajanna L, Systematic Evaluation of Free Radical Scavenging and Antioxidative Activities In *Digera muricata* (L.) Mart., Asian Journal of Pharmacy and Life Science, 1(3), 2011, 249-260.
38. Sharma N, Sharma P and Vijayvergia R, Evaluation of phytochemical and antioxidant activity of some medicinal plants of family Amaranthaceae". Journal of Pharmacy research, 5(9), 2012, 4713-4715.
39. Khan MR, Rizvi W, Khan GN, Khan RA and Shaheen S, Carbon tetrachloride induced nephrotoxicity in rat: Protective role of *Digera muricata*. J. Ethnopharmacol., 122, 2009, 91-99.
40. Hussain A. Evaluation of anthelmintic activity of some ethnobotanicals, Thesis. University of agriculture, Faisalabad, Pakistan, (2008) 87.
41. Weber LW, Boll M, Stampfl A, Hepatotoxicity and mechanism of action of haloalkanes: carbon tetrachloride as a toxicological model, Crit Rev Toxicol., 2003; 33(2), 105-36.
42. Lin HM, Tseng HC, Wang CJ, Lin JJ, Lo CW, Chou FP, Hepatoprotective effects of Solanum nigrum Linn. extract against CCl4-induced oxidative damage in rats. Chemico-Biol. Interact., 171, 2008, 283-293.
43. Reddy KN, Trimurthulu G and Reddy CS, Medicinal plants used by ethnic people of Medak district, Andhra Pradesh, Indian journal of Traditional knowledge, 19, 2010, 184-190.
44. Bindu V and Jain VK, Allelopathic effect of *Digera muricata* (L.) mart on *in vitro* seed germination of *Pennisetum typhoideum*. International journal of plant science. 6(2),2011, 332-334.
45. Khan MR and Younus T, Prevention of ccl4-induced oxidative damage in adrenal gland by *Digera muricata* extract in rat, Pak. J. Pharm. Sci., 24,(4), 2011, 469-473.
46. Khan MR, Memon A, Khan GN, Shabbir M, Saeed N, Shah, Ali, Bokhari Jasia and Rashid U, Protective effects of *Digera muricata* (L.) Mart. against carbon tetrachloride induced oxidative stress in thyroid of rat African Journal of Biotechnology, 10(76), 2011, 17564-17570.

Source of Support: Nil, Conflict of Interest: None.