Abstract: *Tridax procumbens* Linn. (Compositae) is a weed found throughout India. The plant is native of tropical America and naturalized in tropical Africa, Asia, and Australia. Local people known it as “Ghamara”, in English popularly called ‘coat buttons’ and is dispensed for “Bhringraj” by some of the practitioners of Ayurveda. *Tridax procumbens* is a widely occurring medicinal herbs used by Ethnomedical practitioners. It has known for its number of pharmacological activities like Hepatoprotective activity, Anti inflammatory, Wound healing, Antidiabetic activity, Hypotensive effect, Immunomodulating property, bronchial catarrh, dysentery, diarrhoea.

The present study was aimed to investigate phytochemical present in the leaves extract of *Tridax procumbens*. Initially dried powder of *Tridax procumbens* was extracted successively in Chloroform, Acetone-Water and Chloroform-Water and tested for the presence of different phytochemicals.

Keywords: *Tridax procumbens* Linn, Extracts and Phytochemistry.

Introduction

Man and animals depends on the plants for their very existence. Our environment is characterized by richly diversified plant life. Plant diversity is composed of more than 5,00,000 botanical species. Plants constitute a vital component of the biodiversity as they play a key role in maintaining earth’s environmental equilibrium and ecosystem stability. Herbal medicine is known to be the oldest form of healing. It originated from ancient Greek as far back as 1600BC (Baker, H. G. 1970). With Herbal Renaissance happening all over the globe, medicinal herbs are staging a phenomenal comeback. Ethnobotanical information from India estimates that more than 6000 higher plant species forming about 40% of the higher plant diversity are used in its codified and folk healthcare traditions (Ved and Goraya, 2007). In India, Ayurvedic System of medicine has existed for over four thousand years. From ancient literature it is evidence that the various parts of the plants were used in Siddha, Ayurveda and Unani medicines for the treatment of diseases of human being.

*Tridax procumbens* Linn is a common grass found in the tropics. Traditionally, it is
used for the treatment of bronchial catarrh, dysentery, malaria, diarrhea, high blood pressure
and to check haemorrhage from cuts, bruises and wounds and to prevent falling of hair. It
possesses Anti diabetic (Durgacharan et.al 2008), Anti-bacterial (Chitra pai et.al 2011), Anti
plasmodial (Rappiah et.al 2011), Anti hepatotoxic, Anti-oxidant (Reddipalli et.al 2008) and
Antimicrobial (Sneha et.al 2010) properties.

In the present study we investigated 16 phytochemicals qualitatively from various
extracts.

Experimental

Plant Material

The leaves of *Tridax procumbens* Linn. were collected from, Gadhinglaj Tahsil of
Kolhapur district, Maharashtra during Feb 2013. It was authenticated by Prof. R.S. Sawant
Department of Botany, Dr. Ghali College, Gadhinglaj, Kolhapur district, Maharashtra.

Preparation of Extract

Chloroform Extract

The collected leaves of Tridax procumbens Linn were washed and dried under shade. The
coarse powder of the leaves (400 gm) was soaked in 500 ml of Chloroform and extracted in
the cold for 2 days with occasional shaking. The solvent from the total extract was filtered
and filtrate was dried under shade, it was used for phytochemical screening.

Chloroform-Water Extract

The residue obtained from Chloroform extract was mixed with 600 ml of distilled water and
extracted in the cold for 2 days with occasional shaking. The solvent from the total extract
was filtered and filtrate was concentrated on water bath for 3 hrs, it was used for
phytochemical screening.

Acetone-Water Extract

500 gm of coarse powder of leaves of Tridax procumbens Linn were mixed with 300 ml of
distilled water and 300 ml of Acetone in cold for 2 days. The solvent from the total extract
was filtered and filtrate was concentrated on water bath for 3 hrs, remaining filtrate was used
for phytochemical analysis.

Chemicals and Drugs

All the chemicals and solvents were of Analytical grade from SD Fine Chemicals Pvt.
Limited, Bombay.
Phytochemical Analysis

The individual extract was subjected to the qualitative phytochemical screening for the presence of some chemical constituents. Phytochemical test were carried out adopting standards procedure (Trease et. al 1983, Kokate et.al 1997, Hegde et.al 2010). Test were performed for Steroids, Tannin, Saponin, Anthocyanin, Coumarins, Emodins, Alkaloids, Proteins, Amino acids, Diterpenes, Phytosterol, Phenol, Phlobatannins, Leucoanthocyanin, Cardial glycosides and Flavonoids.

Steroid

1ml extract was dissolved in 10 ml of chloroform & equal volume of concentrated H$_2$SO$_4$ acid was added from the side of test tube. The upper layer turns red and H$_2$SO$_4$ layer showed yellow with green fluorescence. This indicates the presence of steroid.

Tannin

a) 2ml extract was added to 1% lead acetate a yellowish precipitate indicates the presence of tannins.

b) 4ml extract was treated with 4 ml FeCl$_3$ formation of green colour indicates that presence of condensed tannin

Saponin

5 ml extract was mixed with 20 ml of distilled water then agitated in graduated cylinder for 15 min formation of foam indicates Saponin.

Anthocyanin

2 ml of aqueous extract is added to 2 ml of 2N HCl & NH$_3$, the appearance of pink red turns blue violet indicates presence of Anthocyanin.

Coumarin

3 ml of 10% NaOH was added to 2 ml of aqueous extract formation of yellow colour indicates coumarins.

Emodins

2 ml of NH$_4$OH and 3 ml of benzene was added to extract appearance of red colour indicates presence of emodins.

Alkaloids

A quantity (3 ml) of concentrated extract was taken into a test tube and 1 ml HCl was added the mixture was heated gently for 20 min cooled and filter, the filtrate was used for following test.
a) Wagner test: Filtrate was treated with Wagner’s reagent; formation of brown reddish precipitate indicates presence of alkaloids.

b) Hager’s test: Filtrate was treated with Hager’s reagent, presence of alkaloids confirmed by the yellow colored precipitate.

Proteins

Xanthoproteic test: Extract was treated with few drops of concentrated HNO\textsubscript{3} formation of yellow indicates the presence of proteins.

Amino acids

Ninhydrin test: To the 2 ml extract 2 ml on ninhydrin reagent was added & boil for few minutes, formation of blue colour indicates the presence of amino acid.

Diterpenes

Copper acetate test: Extract were dissolved in water and treated with 10 drops of copper acetate solution, formation of emerald green colour indicates presence of diterpenes.

Phytosterol

Salkowski’s test: Extract was treated with chloroform and filtered. The filtrate was treated with few drops of concentrated H\textsubscript{2}SO\textsubscript{4} and shakes, allow standing, appearance of golden red indicates the positive test.

Phenol

Ferric Chloride test: Test extract were treated with 4 drops of Alcoholic FeCl\textsubscript{3} solution. Formation of bluish black colour indicate the presence of Phenol

Phlobatannins

Deposition of red ppt when aqueous extract of each plant sample is boiled with 1% Aqueous HCl was taken as evidence for presence of Phlobatannins.

Leucoanthocyanin

5 ml of isoamyl alcohol added to 5 ml of aqueous extract, upper layer appear red in colour indicates the presence of Leuanthocyanin.

Cardial Glycosides

Keller-Killani Test: Plant extract treated with 2 ml glacial acetic acid containing a drop of FeCl\textsubscript{3}. A brown colour ring indicates the presence of positive test.

Flavonoid

a) Alkaline reagent test: Extract was treated with 10 % NaOH solution, formation of intense yellow colour indicates presence of Flavonoid.
b) NH₄OH test: 3 ml of extract were 10 % NH₄OH solution development of yellow fluorescence indicates positive test.

c) Mg turning test: Extract were treated with Mg turning and add conc.HCl to this solution add 5ml of 95 % ethanol, formation of crimson red colour indicates Flavonoid.

d) Zn test: 2 ml extract were treated with Zn dust and conc.HCl development of red colour indicates presence of Flavonoid.

Result and Discussion

Present study deals with qualitative analysis of leaves extract of *Tridax procumbens* Linn. Table no. 1 shows the results of phytochemical analysis of leaves of *Tridax procumbens* Linn. Chloroform extract of leaves of *Tridax procumbens* Linn shows the presence of Steroid, Saponin, Coumarins, Alkaloids, Amino acids, Diterpenes, Phenol and Flavonoids whereas Tannin, Anthocyanin, Emodins, Proteins, Phytosterol, Phlobatannin, Leucoanthocyanin and Cardial Glycosides were absent.

Acetone-Water extract of leaves of *Tridax procumbens* Linn shows the presence of Steroid, Tannin, Saponin, Anthocyanin, Coumarins, Alkaloids, Diterpenes, Phenol and Flavonoids whereas Emodins, Proteins, Amino acids, Phytosterol, Phlobatannin, Leucoanthocyanin and Cardial Glycosides were absent.

Chloroform -Water extract of leaves of *Tridax procumbens* Linn shows the presence of Steroid, Tannin, Saponin, Anthocyanin, Coumarins, Alkaloids, Amino acids, Diterpenes, Phenol and Phlobatannin whereas Emodins, Proteins, Phytosterol, Leucoanthocyanin, Cardial Glycosides and Flavonoids were absent.

Ikewuchi Jude *et al* (2009) was reported Six phytochemical from the leaves of *Tridax procumbens* Linn. Ayyappa Das *et al* (2009) was calculated Eight secondary metabolites from the Aqueous and Methanolic leaf extract of *Tridax procumbens* Linn. Dhanabalans *et al* (2008) also shows the presence of Eight phytochemicals as Alkaloids, Tannin, Saponin, Steroid, Phlobatannin, Terpenoids, Flavonoids and Cardiac glycosides form the Methanolic extract of leaves of *T. procumbens* Linn.
References


Table 1: Phytochemical analysis of leaves of *Tridax procumbens* L.

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<th>C.E.</th>
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+ = Present; - = Absent; A.W.E- Acetone Water Extract; C.E.- Chloroform Extract, C.W.E.-Chloroform Water Extract