Evaluation of Pharmacognostic, Phytochemical and Physicochemical Standards of Wedelia Trilobata (L.) Root

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Abstract

Context: Ethnomedicinally, the root of Wedelia trilobata L. (Asteraceae) has long been used in various ailments in traditional system; most importantly it is used against backache, muscle cramp, rheumatism, stubborn wounds, sores, swelling and arthritic pain, fever and malaria. The main problem experienced in the standardization of herbal drugs is lack of proper identification of plant source. So there is need to establish quality control parameters by using pharmacognostic and phytochemical evaluation, which ensures the purity, safety and efficacy of medicinal plant W. trilobata.

Aim: To evaluate pharmacognostic properties including macroscopic, microscopic and physicochemical parameters of the root of W. trilobata.

Methods: Micro and Macroscopic characters of fresh and dried root samples were investigated. Physicochemical parameters were done by utilizing WHO recommended parameters, preliminary phytochemical and fluorescent analysis of root sample were performed for identification and standardization of root of W. trilobata.

Results: The color, shape, size, odor and surface characteristics were noted from the root and powdered root material of W. trilobata. Light electron microscope images of cross section of root and powdered root revealed that the presence of cork cells, lignified spiral vessels, and parenchymatous cells. Phytochemical screening showed the presence of flavonoids, tannins, phenols, saponins, steroids, carbohydrates and glycosides. Physicochemical parameters such as moisture content, ash value, extractive value and fluorescent behavior of root powder were determined. These parameters are useful tools to differentiate the powdered drug material.

Conclusion: The present study is helpful to supplement the information with regard to its standardization and identification and in carrying out further research in Ayurvedic system of medicine.

Keywords: Pharmacognostic; Microscopical; Wedelia trilobata L; Physicochemical and lignified spiral vessels
Introduction

The procedure of standardization is achieved by pharmacognostic studies that assist in authentication and identification of plant materials. Proper quality and identification assurance of the raw materials are crucial in herbal treatment to insure their quality, safety and efficacy. Pharmacognosy could possibly be a dependable and easy device, by that total information of the crude medication is obtained. The majority of the pharmacopeias and regulatory recommendations are suggested macroscopic and microscopic analysis and chemical substance profiling of natural components for quality control and standardization [1] Wedelia trilobata (L.) Hitchc (Synonym: Sphagneticola trilobata (L.) Pruski), belonging to the Asteraceae family and native to South America, is a perennial creeping herb which is known as an invasive plant at many tropical and subtropical areas including southern China [2,3]. W. trilobata is an herbaceous creeping perennial shrub, up to 70cm height, forms dense mounded mats over the ground. Leaves are glossy green, paler green below, with simple coarse white hairs, serrated margins, sometimes with a pair of lateral lobes.

The stem is rounded, rooting at nodes, stolons up to 2m in length or more. Flowers are borne on solitary or branched inflorescences, ray florets yellow (8-13 per head), central disc florets yellow and tubular, flowers freely produced throughout the year in warm tropics. Fruits are 3-cornered nuts, very small (3 - 5mm), with corky covering and topped by short scales, mature from green to brown, dispersed by water. New plants arise from nodes that root at the soil surface. Seed production is low and generally does not reproduce prolifically via seed [4]. It really is utilized by Indians as traditionally in the treatment of backache, muscle cramp, rheumatism, stubborn wounds, sores, swelling and arthritic pain, fever and malaria [5-7]. So far, phytochemical studies have revealed some structurally diverse chemicals from this plant, including terpenoids (sesqui-, diter-, and triterpenoids), steroids, flavonoids and phenolics, some of which showed significant bioactivities [7-13]. Numerous pharmacological activities of W. trilobata offers been reported such as antimicrobial [14], Antioxidant [15], Anti-inflammatory [15], wound healing [16], Anthelmintic [17] and anticancer [15].

Therefore, we make an effort for standardization of Wedelia trilobata L. root to study the morphological, anatomical, physicochemical and preliminary phytochemical evaluation of root was carried out.

Material and Methods

Plant Material and Authentication

Wedelia trilobata L. Plant was procured in the month of September 2017, from V. V. Institute of Pharmaceutical Sciences, Gudlavalleru, Andhra Pradesh. It had been recognized and authenticated by K. Madhavachetty, plant taxonomist, Division of Botany, Sri Venkateswara University, Tirupati, Andhra Pradesh and voucher specimen of the plant was deposited at the herbarium for long-term reference.

Pharmacognostic Evaluation

Organoleptic evaluation: Organoleptic characteristics of Wedelia trilobata root was assessed by observing color, odor, taste, shape and size according to WHO quality control methods for herbal medicine [18-20].

Microscopic Evaluation

Preparation of sections: Free handed sections of the root were cut into thin sections manually with the sharp cutting edge of the blade. Then transferred on the slide, cleared by warming with chloral hydrate, stained with phloroglucinol and Conc. HCl and mounted in glycerin. The lignified and cellulosic tissues were recognized by utilizing different staining techniques [18].

Powder Microscopy

The powder microscopy was performed according to the method mentioned in Khandelwal [18].

Physicochemical analysis: Physicochemical parameters such as ash value, moisture content and extractive values were determined according to the procedures mentioned in WHO quality control methods for herbal materials [18-20].

Phytochemical analysis: Chloroform, ethyl acetate, ethanol and water extracts of Wedelia trilobata L. were subjected to qualitative chemical analysis of various phytoconstituents like alkaloids, glycosides tannins, flavonoids, steroids and volatile oils according to methods of Khandelwal [18,20-22].

Preparation of extract: The root of Wedelia trilobata was shade dried and powdered. 100g of powdered root was subjected to cold maceration by increasing order of polarity viz chloroform, ethyl acetate, ethanol and water. After 24 hrs filtered the extracts and concentrate with the help of rotary evaporator.

Fluorescence analysis of the powdered drug: The fluorescence examination of the plant material was done by placing the dry powdered root on a slide and observing
by treating with a few drops of different chemical reagents to detect the color changes under UV and visible light [18,23].

**Results**

**Pharmacognostic Evaluation**

**Organoleptic and microscopic evaluation:** The organoleptic characteristics of root showed in Table 1. The transverse section of root is found to be circular in outline. The epidermis is the outer most layer it is made up of cuboidal shaped cells, which are arranges compactly without any intercellular sapaces. The outer layer consists of many uniseriate multicellular hairs. The hypodermal layer is composed of parenchymatous cells with some intercellular spaces. The endodermis showed the presence of phloem and xylem. The phloem is present in between the medullary rays. The medullary rays are parenchymatous and are biserrate in nature. Phloem is well developed and shows the presence of phloem fibres, which are lignified. It also showed the presence of phloem parenchyma. The xylem region was similar to phloem region and was also surrounded by biserrate. Xylem tissue consists of spiral xylem vessels, xylem fibres and xylem parenchyma as shown in Figure 1 to 6.

| Organoleptic characters | Observation Root |
|-------------------------|------------------|
| Colour                  | Buff             |
| Odour                   | Characteristic   |
| Taste                   | No taste         |
| Size                    | 5 to 14 cm       |
| Texture                 | Smooth           |

Table 1: Organoleptic Characteristics of *Wedelia trilobata* L. Root

Figure 1: Morphological Characteristics of *Wedelia trilobata* L. Root.

Figure 2: Transverse section of Root of *Wedelia trilobata* L. Ep: Epidermis; Par: Parenchyma cells; Ph: Phloem; Mx: Meta Xylem; PX: Proto Xylem and XY: Xylem.

Figure 3: Detailed TS of Root showed epidermis and parenchymatous cells. Epi: Epidermis and Par: Parenchyma Cells.

Figure 4: Epidermal cells showed uniseriate multicellular covering trichomes.
Powder microscopy: The powder plant material is buff in color, showed lignified spiral vessels, parenchyma, cork cells, medullary rays and Xylem vessels as shown in Figure 6

Physicochemical evaluation: The various physicochemical parameters of root and root powder i.e., loss on drying, ash value and extractive value were determined. The total ash, acid insoluble ash, water soluble ash, petroleum ether soluble, chloroform soluble, ethyl acetate soluble, alcohol soluble and water-soluble extractive values were shown in Table 2.

| Parameters                                      | Values %w/w |
|-------------------------------------------------|-------------|
| Moisture content (Loss on drying)               | 9.23±0.23   |
| Total ash                                       | 12.25±0.56  |
| Acid-insoluble ash                              | 4.5±1.58    |
| Water soluble ash                               | 2.51±0.86   |
| Petroleum ether soluble extractive value        | 1.82±0.05   |
| Chloroform soluble extractive value             | 3.84±0.44   |
| Ethyl acetate soluble extractive value          | 5.66±0.66   |
| Alcohol soluble extractive value                | 9.85±1.85   |

Table 2: Physicochemical Parameters of root powder of Wedelia trilobata L.

Preliminary phytochemical screening: The preliminary phytochemical screening of the extracts viz., chloroform, ethanol, and water was carried out and the results obtained shown in Table 3.

| Phytoconstituents | Method                      | Aqueous Extract | Ethanol Extract | Chloroform Extract |
|-------------------|-----------------------------|-----------------|-----------------|-------------------|
| Flavonoids        | Shinoda Test                | +               | +               | -                 |
|                   | Zn. Hydrochloride test      | +               | +               | -                 |
|                   | Lead acetate Test           | +               | +               | -                 |
| Volatile oil      | Stain test                  | -               | -               | -                 |
| Alkaloids         | Wagner Test                 | -               | -               | -                 |
|                   | Hager’s Test                | -               | -               | -                 |
| Tannins & Phenols | Fecl₃ Test                  | +               | +               | -                 |
|                   | Potassium dichromate test   | +               | +               | -                 |
| Saponins          | Foaming Test                | +               | +               | -                 |
| Steroids          | Salkowski test              | +               | +               | +                 |
| Carbohydrates     | Molish test                 | +               | +               | -                 |
| Acid compounds    | Litmus test                 | -               | -               | -                 |
| Glycoside         | Keller-Killani Test         | +               | +               | -                 |
| Amino acids       | Ninhydrin test              | -               | -               | -                 |
| Proteins          | Biuret                       | -               | -               | -                 |

*“+” Present and “-” Absent

Table 3: Phytochemical analysis of various extracts of Wedelia trilobata L. Root
Fluorescence analysis: The behavioral changes of the powdered drug with distinctive chemical reagents were determined at both UV and Visible light and it is reported as shown in Table 4.

| Solvent used            | Visible light | UV light At short (254nm) | At Long (366nm) |
|-------------------------|--------------|---------------------------|-----------------|
| Distilled water         | Buff         | Buff                      | Dark Brown      |
| 1 N NaOH 1 N Methanol   | Brown        | Black                     | Black           |
| 1 N HCl                 | Buff         | Reddish brown             | Black           |
| 50% HNO₃                | Brown        | Brownish white            | Black           |
| FeCl₃                   | Buff         | Yellowish brown           | Bluish red      |
| CHCl₃                   | Buff         | Yellowish grey            | Dark brown      |
| Picric acid             | Brownish yellow | Yellowish brown    | Black           |

Table 4: Fluorescence analysis of Wedelia trilobata L. Root powder.

Discussion

To assure the reproducible quality of herbal drugs, proper control of starting material is vital. The primary step towards ensuring starting material is authentication. Thus, in recent years there has been a rapid increase in the standardization of medicinal plants. Though modern techniques are available, but still identification of medicinal plants is more reliable on pharmacognostic studies [24]. In this regard, the macroscopic and microscopic features of root have been studied. Macroscopical characters of the root of the plant can serve as a diagnostic parameter. Microscopical study and powder analysis of the plant sample revealed the presence of cork cells, lignified spiral vessels and parenchymatous cells. Further, this study can also be useful to reduce the possibilities of adulteration of this useful herbal drug when it is available in the powdered form [25]. Studies of physicochemical parameters can serve as an important source to judge the purity and quality of crude drugs. The extractive values give the approximate measure of their chemical constituents and from the study, the extractive values of water were highest followed by alcohol. The ash value represents the earthy matter or inorganic components and other impurities present along with the herbal drug. The pharmacognostic standard for the root of Wedelia trilobata L. laid down for the first time in the study. The phytochemical investigation of different solvent extracts viz., chloroform, ethanol and water were examined and it revealed the presence of flavonoids, tannins, phenols, saponins, steroids, carbohydrates and glycosides.

Conclusion

The data generated from the current study facilitate to authenticate the medicinally important plant W. trilobata. Microscopic features could also be useful for establishing the pharmacopeia standards. Morphology as well as various pharmacognostic aspects of the root of W. trilobata was studied and described along with phytochemical and physicochemical parameters that can be useful in further isolation and purification of medicinally important compounds.

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