

RESEARCH ARTICLE

PHARMACOGNOSTIC EVALUATION OF GOMPHRENA SERRATA ROOT

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Article Info:

Abstract



Cite this article:

Article History: Received: 3 June 2017 Reviewed: 6 July 2017 Accepted: 27 August 2017

Published: 15 September 2017

Prasanth DSNBK, Prasanna MM, Priyanka M, Pala NN, Lakshmi PB, Mounika Y, Rao AL. Pharmacognostic Evaluation of *Gomphrena serrata* Root. Universal Journal of Pharmaceutical Research. 2017; 2(4): 6-10. http://doi.org/10.22270/ujpr.v2i4.R2

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DSNBK Prasanth, Department of Pharmacognosy, V.V. Institute of Pharmaceutical Sciences, Gudlavalleru, Andhra Pradesh, India; Tel: +91-7382027437. E-mail: *dsnbkprasanth@gmail.com* **Objectives:** *Gomphrena serrata* (Amaranthaceae) has been utilized for many ailments in the conventional system ethnomedicinally; most significantly against bronchial asthma, diarrhea, hay fever, pains, tonic, carminative, diabetes, dermatitis, and piles. The key challenge experienced in the standardization of herbal drugs is the correct identification of the plant source. Thus, setting up quality control parameters by means of pharmacognostic and phytochemical analysis which assures the purity, safety, and efficiency of *G. serrata* is necessary. The current research was conducted to assess the pharmacognostic characteristics including macroscopic, microscopic, phytochemical and physicochemical parameters of the root of *G. serrata*.

Methods: Micro, as well as macroscopic characteristics was investigated. Physicochemical parameters had been done by implementing WHO suggested parameters; preliminary phytochemical and fluorescent evaluation of root was executed for appropriate identification and standardization.

Results: The color, shape, size, odor, and surface characteristics were reported from the root and powdered root material of *G. serrata*. Light microscope images of cross section and powdered root revealed the presence of lignified xylem fibers, xylem vessels, cork cells and parenchyma cells. Phytochemical testing confirmed the presence of alkaloids, carbohydrates, saponins, tannins, proteins, amino acids, phytosterols and flavonoids. Physicochemical parameters such as moisture content, ash value, extractive value and fluorescent behavior of root powder have also been established

Conclusion: The current research is useful in order to supplement the information with regard to its standardization, identification and in carrying out further investigation in Ayurvedic system of medicine.

Keywords: Gomphrena serrata, lignified xylem vessels, pharmacognostic, phytochemical analysis.

INTRODUCTION

Medicinal plants tend to be playing a crucial role in conventional medicines for remedy of different health problems. On the other hand, a vital barrier, that has obstructed the promotion in the utilization of alternative medicines in the developed nations, is no proof of documentation and lack of stringent quality control measures¹. There is also a requirement for the records of all the research work meted out on conventional medicines by means of documentation. With this particular problem, it has become essential to make assurance regarding the standardization of the plant and its parts to be utilized as a medicine. In the process of standardization, we can make use of various techniques and methodology to attain our objective in a stepwise manner e.g. pharmacognostic and

phytochemical studies. These methods and procedures are useful in identification and standardization of the plant material. Proper characterization and quality assurance of beginning material is an important step to make sure reproducible quality of herbal medicine to help us to rationalize its safety and efficacy. For this reason, we have carried out pharmacognostic studies of *Gomphrena serrata* belongs to family Amaranthaceae¹. This kind of study will not only assist in authentication but also assures reproducibility of herbal products in marketing².

In the current study, investigation was emphasized on one of the commonly available plant in India *i.e.*, *G. serrata*, belongs to family Amaranthaceae. The family Amaranthaceae contains nearly 60-70 exotic species. The genus *Gomphrena*, with around 138 species, some of the important species include *G. boliviana*. *G.* celosioides, G. globosa, G. haenkeana, G. macrocephala, G. martiana, G. meyeniana, G. perennis and G. pulchella³. All parts of this plant are widely used as a folklore medicine for the treatment of various ailments by the Indian traditional healer. Traditionally, the plant is utilized in the remedy of bronchial asthma, diarrhea, hay fever, pains, tonic, carminative, diabetes, dermatitis and piles⁴⁻⁷. G. serrata is annual, ascending or erect herbs, up to 40 cm tall; branches clothed with white, shaggy hairs; leaves are obovate-lanceolate, 2-4X1-1.5 cm, glabrescent above, long white shaggy hair below, obtusely apiculate and the base is cuneate; flowers are white with yellow tinge in axillary and terminal compressed, cylindrical spikes; utricles enclosed hardened perianth; seeds are brown and shiny⁸.

Phytochemical constituents have been separated from the genus Gomphrena i.e., oleuropein³, stigmasterol, βsitosterol, isochavicinic acid, campesterol, betalain, friedelin, 3-epi-friedelinol, allantoin, and chrysoeriol- $7-O-\beta-D-glucoside^9$. Ethnomedicinally, the genus Gomphrena has been documented various pharmacological activities including antimicrobial¹⁰, anticancer¹¹, antimalarial¹², and analgesic¹³. Although the plant has been extensively used for its traditional value, pharmacognostic, phytochemical and pharmacological account remains unexplored. Therefore the current investigation had been carried out to study the morphological, microscopical, physicochemical and phytochemical characteristics of the root of G. serrata with the purpose of contributing to the establishment of monograph^{14,15}.

MATERIALS AND METHODS

Plant collection and authentication

The plant obtained from Tirupati, Chittoor district of Andhra Pradesh, India during the month of December 2016 and authenticated by Dr. K. Madhava chetty, Taxonomist at Sri Venkateswara University Tirupati, India. Voucher specimen No. 1864 was deposited at the herbarium for future reference. One portion of the root is preserved in formalin: acetic acid: alcohol mixture for histological studies and the remaining portion was shade dried, powdered and sieved through 20 mesh and kept in an air tight container for future use.

Chemicals

All analytical grade chemicals i.e., absolute alcohol, phloroglucinol, acetic acid, chloral hydrate, H₂SO₄, NaOH, HNO₃, FeCl₃, conc. HCl and chloroform were utilized in this study, which was procured from E. Merck, Germany.

Pharmacognostic evaluation

Morphological evaluation

Organoleptic evaluation of *G. serrata* root has been carried out in accordance the color, size, odor, shape, and taste as per WHO Quality Control methods of herbal medicine¹⁶.

Microscopic evaluation

Preparation of sections

Microscopic studies had been done by preparing thin hand section of the root with the help of sharp cutting edge of the blade, then cleared with chloral hydrate solution, stained with phloroglucinol-hydrochloric acid (1:1) and mounted in glycerin.

Powdered microscopy

The powder microscopy was carried out in accordance with the procedure described in Khandelwal¹⁷.

Preparation of extracts and preliminary phytochemical analysis

The powdered material had been extracted with various solvents according to its polarity i.e., chloroform, methanol, and water. Five grams of root powder were extracted with 20 ml of the respective solvent by maceration at room temperature for 24 hours. Then, filtered through Whatman filter paper and collect the filtrate, concentrated with roto-evaporator. Then, the extracts had been subjected to preliminary phytochemical screening according to standard methods^{17,18}.

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 materials¹⁶.

Fluorescence analysis

Various reagents were utilized to check the fluorescence activity. In this, 0.1 g of root powder was blended with 1.5 ml of respective reagent (Table 4). The mixture was placed on a slide for a minute and observed under visible light, short ultra-violet light (254 nm) and long ultraviolet light (365 nm)¹⁹.

RESULTS

Morphological characteristics

The morphological characteristics of *G. serrata* root were described in Figure 1 and Table 1.



Figure 1: Organoleptic characteristics of the whole Plant of *G. serrata*.

Anatomical description Root

The transverse section of the root of *G. serrata* showed the presence of Cortex was made up of thin walled parenchymatous cells with very small intercellular spaces. Cork showed the presence of periderm i.e., 2-3 layered narrow, tangentially elongated cells with dark brown granular matter. Phelloderm is 1-2 layered rows of tangentially elongated thin walled cells. 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 uniseriate to triseriate, majorly biseriate. Radially arranged vascular bundles were present in which, Phloem is well developed and shows the presence of phloem fibers, which are non-lignified. It also showed the presence of phloem parenchyma.

Table 1: Morphological characteristics of root of G. serrata.

Characters	Observation		
Colour	Buff		
Odour	Characteristic		
Taste	Characteristic		
Texture	Smooth		
Thickness	4-12 cm		

The xylem region was similar to phloem region and was also surrounded by uniseriate to triseriate medullary rays. Xylem tissue consists of spiral xylem vessels, xylem fibers and xylem parenchyma (Figure 2).



Figure 2: Transverse section of the root of *G*. serrata. Ck: Cork; Par: Parenchyma; Xy: Xylem; Ph: Phloem; Px: Proto Xylem and Mx: Meta Xylem.

Phytoconstituents	Method	Aqueous	Methanolic	Chloroform	Pet. ether
-		extract	extract	extract	extract
Flavonoids	Shinoda Test	+	+	-	-
	Zn. Hydrochloride test	+	+	-	-
	Lead acetate Test	+	+	-	-
Volatile oil	Stain test	-	-	-	-
Alkaloids	Wagner Test	+	+	+	-
	Hager's Test	+	+	+	-
Tannins	FeCl ₃ Test	+	+	-	-
and Phenols	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	+	+	-	-
	+ Pr	esent - Abser	nt		

Table 2. Preliminary qualitative phytochemical analy	sis of G serrata root

Powder microscopy

The crude powder of root was buff in color with characteristic odor and taste. Microscopic study of the powder showed revealed different characters such as cork cells, parenchyma cells, lignified xylem vessels, and xylem fibers (Figure 3).



Figure 3: Powder microscopy of root of G. serrata.

Preliminary qualitative phytochemical analysis The results of qualitative phytochemical analysis of crude powder of G. serrata root are shown in Table 2.

Physicochemical parameters

The results attained from various determinations of physicochemical analysis are produced in Table 3.

Fluorescence analysis

Fluorescence analysis of root powder was performed out after treating with different solvents. Fluorescence was observed at 254 nm and 365 nm comparing its change of color in the visible light. The observations presented in Table 4 show the variation in color.

Table 3: Physicochemical parameters of root powder of G. serrata.

Parameters	Values	
	%w/w	
Moisture content (Loss on drying)	8.52±0.53	
Total ash	4.86 ± 0.45	
Acid insoluble ash	3.25 ± 0.18	
Water soluble ash	2.22 ± 0.47	
Petroleum ether soluble extractive	0.23 ± 0.08	
value		
Chloroform soluble extractive value	1.88 ± 0.04	
Ethyl acetate soluble extractive	3.56 ± 0.04	
value		
Methanol soluble extractive value	5.98±0.12	
Water soluble extractive value	7.62 ± 0.08	

Solvent used	Visible light	UV light		
		At short (254 nm)	At long (365 nm)	
Distilled water	Buff	Dark Brown	Buff	
Methanol	Brown	Black	Greyish green	
1N HCl	Buff	Dark Brown	Green	
50% HNO3	Buff	Brown	Yellowish Black	
FeCl ₃	Orange	Dark blue	Black	
CHCl ₃	Pale green	Buff	Black	
Picric acid	Yellowish buff	Dark blue	Fluorescent blue	
Ethyl acetate	Buff	Buff	Greenish black	

Table 4: F	luorescence a	nalysis of (G. serrata	root powder.
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DISCUSSION

Indian systems of medicine utilize the majority of the crude drugs which are of plant origin. It is important that standards need to be set down to control and check the identity of the plant and confirm its quality before use. Hence a detailed pharmacognostic assessment is an extremely an important prerequisite. In accordance with World Health Organization (WHO), the organoleptic and histological description of a medicinal plant could be the first step towards establishing its identity and purity and should be performed before to any tests tend to be undertaken²⁰. G. serrata, extensively utilized in conventional medicines has tremendous therapeutically potential due to its various biological activities. The prominent diagnostic characteristics of the root were xylem fibers, lignified xylem vessels, cork cells and parenchymatous cells. These characters can be utilized for standardization of drugs as well as used for preparation of plant monograph and also reduces the possibilities of adulteration, when the drug is available in the powdered form studies of physicochemical parameters can serve as an important source to judge the purity and quality of crude drugs. Ash values are utilized to establish the quality and purity of the crude drug. It implies the existence of various impurities like carbonate, oxalate, and silicate. The water soluble ash is water soluble part of total ash, employed to calculate the amount of inorganic substances found in the drugs. The acid insoluble ash comprises mostly silica and indicates contamination with earthy matter. The moisture content of drugs might be at the minimum level in order to suppress the growth of microorganisms like bacteria, yeast or fungi during storage. The extractive values are helpful to judge the chemical constituents present in the crude drug and also assist in the evaluation of particular constituents soluble in a specific solvent. Total ash and acid insoluble ash are essential indices that illustrate the quality and purity of the herbal medicine. Total ash consists of physiological ash, which is derived from plant tissue itself, and non physiological ash that is usually derived from atmosphere contaminations including sand and soil. Total ash content alone is not adequate to indicate the quality of herbal medicine because the plant materials usually contain a significant level of physiological ash, calcium oxalate in particular. Therefore, the acid insoluble ash content is another index to indicate the quality of herbal medicine. The phytochemical analysis of extracts viz.,

petroleum ether, chloroform, methanol, and water was analyzed and it indicated the presence of alkaloids, carbohydrates, saponins, tannins, proteins, amino acids, phytosterols, and flavonoids²¹⁻²³.

CONCLUSIONS

Standardization of herbal drugs is very much crucial because they are produced from heterogeneous sources which could result in variations. These kinds of variations can cause spurious results in various pharmacological and phytochemical studies. *G. serrata* root was recognized for many therapeutical properties, therefore, the current study might be beneficial to supplement the information in respect to its identification, authentication, and standardization; no such information is available for the same till date.

AUTHOR'S CONTRIBUTION

DSNBK: Prasanth writing original draft, methodology, investigation, formal analysis, conceptualization. Prasanna MM: writing, review and editing. methodology, formal analysis, conceptualization. Priyanka M: writing, review, and editing, methodology. Pala NN: writing, review, and editing, Lakshmi PB: writing, review and editing. Mounika Y: formal analysis, writing, review, and editing. Rao AL: writing, review, and editing, investigation, conceptualization. All authors revised the article and approved the final version.

ACKNOWLEDGEMENTS

The authors extend their thanks and appreciation to the Institute of Pharmaceutical Sciences, Gudlavalleru, Andhra Pradesh, India to provide necessary facilities for this work.

DATA AVAILABILITY

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

CONFLICT OF INTEREST

None to declare.

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