# Macro-microscopic and physico-chemical details of *Rostellularia procumbens* (L.) Nees var. simplex (D.Don) Yamasaki

B. JYOTHI, BULUSU SITARAM, G. SUDARSANAM, M. NAGALAKASHMI DEVAMMA, CH. PARAMAGEETHAM AND G. PRASADA BABU

Accepted: November, 2009

## **SUMMARY**

Rostellularia procumbens var. simplex is used in several Ayurvedic drugs for the treatment of asthma, cough, bone diseases and rheumatism. The botanical, macro-, microscopic characters, macerate, powder analysis, histochemical tests and physicochemical details are presented.

**Key words:** Macro-microscopic characters, Macerate, Physical constants fluorescence studies.

The genus *Rostellularia* is distributed in India and Sri Lanka. The genus consists of 21 species (Gamble and Fischer, 1967). The genus Rostellularia procumbens is being pharmacologically and chemically screened in recent years (Mruthyunjaya Swamy et al., 1998; Chen et al., 1998; Weng et al., 2004). However in R.procumbens var.simplex pharmacognostical work is lacking. Some of the medicinal properties attributed to Rostellularia procumbens var. simplex are whole plant is used as medicine in asthma, cough, rheumatism, backache, flatulence and lumbago. Decoction of the leaves is sometimes used in the diseases of the bone (Anonymous, 1959). A perusal of the literature revealed that no pharmacognostical work has been carried out (Gurudeva and Yoganarasimhan 2009). Hence, the present work to cover the morphology of the taxon, macro-, microscopic characters of the root, stem and leaves, physical constants and fluorescence were studied.

## MATERIALS AND METHODS

The plant material was collected from Japalitheertham, Tirumala hills and the voucher herbarium specimen was processed followed by standard methods (Jain and Rao, 1977). Macro and microscopical studies were carried out (Johansen, 1940 and Wallis, 1985) during the year 2005.

## Correspondence to:

**B. JYOTHI**, Department of Botany, S.P.W.D. and P.G. College, TIRUPATI (A.P.) INDIA

Authors' affiliations:

**BULUSU SITARAM**, Department of Dravyaguna, S.V. Ayurvedi College, TIRUPATI (A.P.) INDIA

G. SUDARSANAM, M. NAGALAKASHMI DEVAMMA AND G. PRASADA BABU, Department of Botany, Sri Venkatswara University, TIRUPATI (A.P.) INDIA

**CH. PARAMAGEETHAM**, Department of Microbiology, Sri Venkatswara University, P.G. Centre, TIRUPATI (A.P.) INDIA

Physical constants were carried out by standard methods (Kokoski *et al.*, 1958; Chase and Pratt, 1949; Krebs *et al.*, 1969) and fluorescence studies followed by standard procedures (Khandelwal *et al.*, 1996).

# Taxonomy:

Diffuse herb, branchlets strigose-hispid, upto 30 cm. Leaves elliptic, 4-4.5 x 1.5 cm scabrous above, glabrescent to strigose below, acute at both ends, petiole upto 0.5 cm spikes terminal, 1-2 cm, bracts obovate, 4 x2 mm, scarious, midrib broad, shortly excurrent, margin ciliate acute, bracteoles lanceolate, 4x0.7 cm, midrib, broad, margin thinly scarious, ciliate, acute. Calyx lobes-4, some times with a 5th rudimentary lobe, sub-equal, lanceolate 4 mm, midrib ridged, scarious, closely ciliate, acute. Corolla 6x3 mm across, tube 3.5 mm, upper lip 3.5 mm, truncate shortly, emarginate, lower lip sub orbicular, 5x5 mm puberulous without. Stamens 2 to 3 mm anthers, 0.7 mm. Ovary 1.5 mm, pubescent, style 4 mm, capsule oblong, 4 x1.5 mm sparsely hairy, pointed, seeds 1 mm across (Plate:1)

## Herbarium specimen examined:

Author (212) collected on 23<sup>rd</sup> November 2006, on way to Japalitheertham, Tirumala hills, Chittoor district of Andhra Pradesh and it is authenticated with D.Rangacharyulu(2292) deposited at the Herbarium of S.V.University, Tirupati.

# RESULTS AND DISCUSSION

The results obtained from the present investigation are presented below:

#### Root:

*Macroscopical characters:* 

Fairly thick, 1.1 mm thickness, tap root is small,

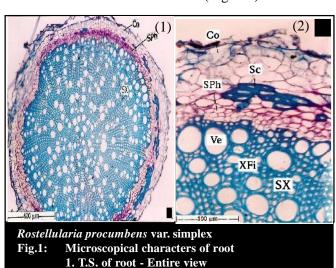


lateral roots are very slender, hairy, bitter and no specific odour.

(D.Don) Yamasaki (Acanthaceae)

# Microscopical characters:

Transverse section of the root is circular in outline and shows epidermis, cortex and vascular cylinder. Epidermis of the root is broken and periderm is formed in the region where epidermis is removed. Cortex is narrow having 2 to 3 layers of tangentially stretched parenchyma. Discontinuous patches of sclereids are observed inner to the cortex. Secondary phloem occurs all around the xylem cylinder. Secondary xylem is solid and dense with uneven outline (Fig. 1.1). It consists of wide vessels thick walled and libriform fibres. Vessels are diffuse in distribution and are mostly solitary, include both wide and narrow vessels. Widest vessel is 30 mm in diameter, the narrow vessel is 15 mm diameter (Fig. 1.2).



[Internat. J. Plant Sci., Jan. - June, 2010, 5 (1)]

2. T.S. of root – A sector enlarged

#### Stem:

Macroscopical characters: Stems are branched, woody, 2 cm thickness, easily peelable and slightly bitter to taste.

# Microscopical characters:

Transverse section of the stem shows wide ridges and shallow furrows (Fig. 2). It consists of epidermis, cortex and wide stelar region. Epidermis is thin with narrow tabular cells in the furrow region, in the ridged region epidermal cells are dilated and radially elongated. 1-2 layers of colleen-chymatous hypodermis is present beneath the epidermis of the ridges. In furrows the cortex is made up of three layers of chlorenchyma.

A distinct endodermis is present around the vascular cylinder, it consists of barrel shaped thin walled cells. Secondary phloem is narrow and made up of 4 layers of cells. Secondary xylem is prominent and continuous. It consists of alternating radial bands of vessels and fibres. Vessels are circular, narrow thin walled, solitary and diffuse in distribution. Fibres are in regular radial rows, they are thick walled and lignified. Xylem cylinder is 170 mm thick. Primary xylem occurs in ten or more points

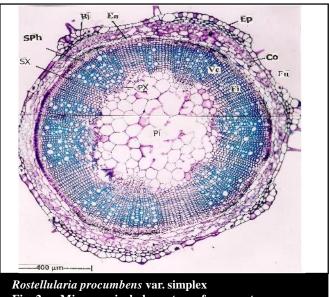


Fig. 2: Microscopical characters of young stem

around the pith. Central portion is occupied by wide pith cavity due to disintegration of pith cells.

## Leaf:

Macroscopical characters: Leaves elliptic, 4 - 4.5 x 1.5 cm across, acute at both ends and taste bitter.

# Microscopical characters

Transverse section of the leaf shows epidermis, mesophyll and vascular bundle. Leaf has prominent lateral vein (Fig. 3.1) and midrib (Fig. 3.2) with uniformly thin lamina prominent conical adaxial cone and semicircular abaxial part (Fig. 3.1). Midrib has slightly broad adaxial hump and flat rectangular abaxial part. Epidermis is made up of squarish cells with thin cell walls. A small group of 2 to 3 layers of collenchyma cells are present in midrib zone just beneath the epidermis. Palisade parenchyma cells are horizontally transcurrent across the adaxial part. Abaxial midrib has prominent layer of epidermis with thick walls and prominent cuticle. Mesophyll made up of thin walled, compactly arranged parenchyma. Lateral vein is 350 mm in vertical plane and 200 mm in horizontal plane. Midrib is 450 mm and 350 mm horizontally. Vascular bundle of the lateral vein is single top-shaped and collateral (Fig. 3.1). The vascular bundle of the midrib is slightly broader and bundle sheath is absent.

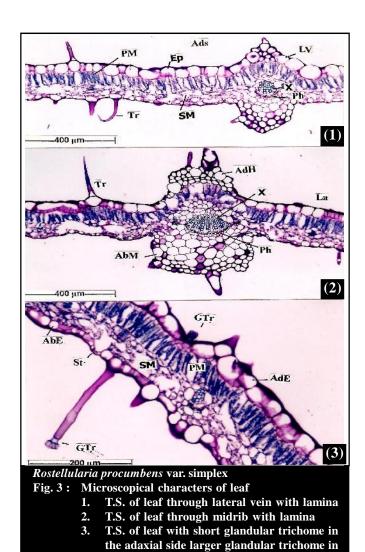
Lamina is 150 mm thick it has thick adaxial epidermis with large squarish cells which are 30-40 mm thick. Abaxial epidermis is thin walled and stomatiferous. Mesophyll consists of single layer of thin walled, loosely arranged palisade parenchyma and 4 to 5 layers of spongy parenchyma with wide air cavities (Fig. 3.1-2). Palisade zone is 80 mm in length.

# Epidermal trichomes

Two types of trichomes are present on the lamina and veins.

Nonglandular or covering trichomes:

This type of trichomes are more abundant. It is two or three celled, uniseriate, unbranched and thin walled pointed tip (Fig. 3.1-2). Trichome may be straight or curved. Basal epidermal cell of trichome is dilated. Trichome is about 220 mm long.



AdE PM Rostellularia procumbens var. simplex Fig. 4: Distribution pattern of cystolith and stomatal morphology T.S. of lamina showing cystoliths Paradermal section showing cystolith distribution Abaxial epidermis with stomata

the abaxial side
[Internat. J. Plant Sci., Jan. - June, 2010, 5 (1)]

#### Glandular trichomes:

These are two types. One type of trichome is small spherical type situated in shallow cavity of the epidermis, it has a short stalk and globular multicellular body with dense content (Fig. 3.3). Second type of trichome has long, unicellular, uniformly thick short stalk cell and a crown of rosette cells (Fig. 3.3). This type is less frequent and occurs mostly on the lower epidermis. Trichome is 450 μm long, terminal rosette of cells 30 μm wide.

## Cystoliths:

Calcium carbonate cystoliths are abundant in the epidermal cells of leaf and young stem. Cystoliths are long and cylindrical with warty surface. They occur in the enlarged epidermal cells called lithocysts. Cystoliths are 220  $\mu$ m long and 20  $\mu$ m width (Fig. 4.1-2).

#### Stomata:

Stomata occur only in lower epidermis of lamina and on young stem (Fig. 4.3). Stomata slightly raised above the epidermal level. Stomata are diacytic. Most of the stomata have more than one circle of subsidiary cells. The stomata are  $11 \mu m^2$ . The guard cells are  $90 \times 50 \text{ mm}$  size.

## Whole plant-macerate:

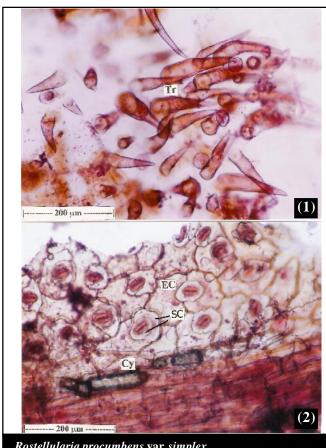
Maceration of whole plant shows the following elements:

# Epidermal trichomes (Fig. 5.1):

Trichomes are one to three celled, uniseriate, unbranched, broad at the base, narrow and conical at the tips, walls thin and smooth, 150 - 180 µm long - 40µm broad at the base and 10  $\mu m$  broad at the apex.

# Vessel elements (Fig. 5.2):

Long, narrow and cylindrical, performation plate simple, horizontal, tails absent, lateral wall pits circular, dense and alternate, length of the vessel elements 240- $340 \mu m$ , width  $20-25 \mu m$ .



Rostellularia procumbens var. simplex Whole plant - Macerate

- Narrow and wide fibres
- Two long cylindrical vessel elements

NFi - Narrow fibers; WFi - Wide fibers;

VE - Vessel elements; PP - Perforation plate



Rostellularia procumbens var. simplex

- Fig. 6: Whole plant Macerate
  - **Epidermal trichomes**
  - Abaxial epidermis with diacytic stomata - Trichome; EC - Epidermal cells;
    - SC Subsidiary cells; Cy- Cystoliths

# Fibres (Fig. 5.1; 6.2):

Both wide and narrow fibres observed. Wide fibres thin walled with wide lumen, 320  $\mu$ m long and 15  $\mu$ m wide (Fig. 5.1). Narrow fibres thick walled with narrow lumen, long with gradually tapering ends (Fig.6.2), 450  $\mu$ m long and 10  $\mu$ m wide. Pits not evident both in the narrow and wide fibres.

# Stomata (Fig. 6.2):

Stomatal type diacytic, epidermal cells slighty lobed, thin walled, guard cells 30x20 µm in size.

# Cystoliths (Fig.7.1,2):

Elongated, cylindrical, either straight or curved, occur in lithocyst, cystolith surface minutely spiny and 150-400  $\mu m$  x 40  $\mu m$  in size.

# Powder analysis:

The powder is green in colour. It is bitter to taste. The observations are given in Table 1.



[Internat. J. Plant Sci., Jan. - June, 2010, 5 (1)]

Cy - Cystolith; Li - Lithocyst

Table 1 : Powder analysis	
Treatments	Observation
Powder treated with water	Non-sticky
Powder shaken with water	Foam like froth
Powder treated with 5% aqueous NaOH	Green
Powder treated with 60% aqueous sulphuric acid	Dark green
Powder pressed between filter paper for 24 hours	No oil stain

#### Histochemical tests:

The sections were treated with different reagents and the observations are provided in Table 2.

Table 2 : Histochemical tests					
Drug	Reagents	Test for	Reaction	Results	
Section	Iodine solution	Starch	Blue colour	+	
Section	Ferric chloride	Tannin	Black	+	
	solution				
Section	Sudan III solution	Oil	No	-	
		globules	Effervescence		
Section	Phloroglucinol +	Lignin	Magenta	+	
	dil. HCl + Alcohol				
Section	Conc. HCl	Crystals	No	+	
			Effervescence		

+ = Present ; - = Absent

# Physico-chemical constants:

Physical constants:

The physical constants determined by standard methods are given in Table 3.

Table 3 : Physical constants	
Total ash%	15.47
Water soluble ash%	4.79
Alkanity of water soluble ash%	5.26
Acid in soluble ash (%)	1.14
Extractive values (%)	
a) Alcohol soluble extract	1.05
b) Water soluble extract	2.09
c) Hexane soluble extract	0.44
d) Chloroform soluble extract	0.92

## Fluorescence analysis:

Fluorescence analysis was carried out by standard procedures. The results are given in Table 4.

#### Conclusions:

It is evaluated for its identification, botanical, macro-

Table 4 : Fluorescence analysis					
Experiments	Visible / day light	UV light			
Experiments	Visible / day light	254 nm	365 nm		
Drug powder	Pale green	Green	Brown		
Drug powder + 1 N NaOH (aq.)	Green	Yellowish green	Black		
Drug powder + 1N NaOH (alc.)	Green	Yellowish green	Green		
Drug powder + 1 N HCl	Pale green	Pale green	Black		
Drug powder + 50% H <sub>2</sub> SO <sub>4</sub>	Dark green	Yellowish green	Yellowish green		
Drug powder + 50% HNO <sub>3</sub>	Orange	Green	Black		
Drug powder + Picric acid	Green	Yellowish green	Yellowish green		
Drug powder + Acetic acid	Green	Green	Black		
Drug powder + Ferric chloride	Dark olive (green)	Pale green	Black		
Drug powder + HNO <sub>3</sub> + NH <sub>3</sub>	Reddish orange precipitate	Yellowish green	Green		

, microscopic and physico-chemical details. Whole plant possess medicinal properties. It is not only used in several Ayurvedic preparations but also used in the preparations of Siddha system of medicine.

## Acknowledgement:

The authors are grateful to the PARC, Chennai, for providing the necessary facilities to carry out this work.

# REFERENCES

- Anonymous, (1959). The Wealth of India (Raw materials), C.S.I.R, New Delhi, Vol. V, 16.
- Chase, C.R. and Pratt, R.J. (1949). Fluorescence of powdered vegetable drugs with particular resource to development of a system of identification. *J. American Pharm. Asso.*, **38**: 324-333.
- Chen, C.C., Hsin, W.C. and Huang, Y.L. (1998). Six new-diarylbutane lignans from *Justicia procumbens*. *J. Natural products.*, **61**(2): 227-229.
- Gamble, J.S. and Fischer, C.E.C. (1967). Flora Of The Presidency of Madras, Vol. 1-3, Calcutta.
- Gurudeva, M.R. and Yoganarasimhan, S.N. (2009). Bibliography Of Medicinal Plants of India. Divyachandra Prakashana Publishers, Bangalore.
- Jain, S.K. and Rao, R.R. (1977). *Field And Herberiam Methods*. Today and Tomorrow publishers, New Delhi.
- Johansen, D.A. (1940). *Plant microtechnique*. McGrow Hill New York, 183-203.
- Khandelwal, K.R., Pawar, A.P., Kokate, C.K. and Gokhale, S.B. (1996). *Practical Pharmacognosy Techniques and Experiments*, Nirali Prakasam, 3<sup>rd</sup> Edition, 140-141.

- Kokoski, C.J., Kokoski, R.J. and Salma, F.J. (1958). Fluorescence of powdered vegetable drugs under ultra-violet radiation. *J. American Pharm. Asson.*, **47**:715.
- Krebs, K.G., Heunsen, D. and Wimmer, H.W. (1969). IN (editor E. Stahl), *Thin Layer Chromatography A Laboratory Handbook* (Second Ed.) London, 204-255, 855-909.
- Mruthyunjaya swamy, B.H.M., Rudresh, K., Swamy, H.K., Badami, S.M. and Hiremath, S. (1998). Antiinflammatory activity of alcohol extract of *Justicia procumbens* (Acanthaceae). *Indian J. Pharma. Sci.* **60** (3): 173-175.
- Rangacharyulu, D. (1991). Floristic studies of Chittoor district, Ph.D. Thesis, S.V. University, Tirupati.
- Wallis, T.E. (1985). *Text Book of Pharmacognosy* (15<sup>th</sup> Ed.), T.A. Churchill, London, 571-582.
- Weng, J.R., Ko, H.H., Yeh, T.L., Lin, H.C. and Lin, C.N. (2004). Two new arylnaphthalide lignans and antiplatelet constituents from *Justicia procumbens Arch. Pharm* (Weinheim), **337** (4): 207-12.

\*\*\*\*\*\*