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Identity parameters on traditionally used Antiurolithiatic Herb - *Scoparia Dulcis* Linn.

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ABSTRACT

Introduction: *Scoparia dulcis* Linn. locally known as *Manithumbe Gida* belongs to *Scophularaceae* family and used in medicine by the traditional practitioners for the treatment of urinary calculi.

Materials and Methods: Matured plants are collected from Udupi district and authenticated. Macro-microscopic features, physico-chemical standards, HPTLC and secondary metabolites were recorded as per standard guidelines. **Result:** TS of leaf has shown the presence of mesophyll and bi-collateral vascular bundles. Outer cork tissue, a layer of cortex and conjoint collateral closed vascular bundles and central pith are inclusions of stem TS. Pitted and reticulate vessels are characteristic features of plant powder. Physico-chemical standards and presence of alkaloids, carbohydrates, tannin and coumarins were indicative of its chemical nature. HPTLC fingerprints are a record of its different chemical constituents. Thus the quality monograph prepared on this drug beneficial in future research.

Key words: *Scoparia dulcis*, Quality standards, HPTLC, Monograph.

INTRODUCTION

Plants are a source of food, cosmetics, medicine and many more since centuries. Though there exists innumerable number of these green sources, only few are identified, used and popularized.^[1] Urbanization and simultaneous growing demand on herbal products are measure zone where one has to open his eyes towards all natural sources of drugs. It is an accepted fact that in principle all natural resources are having medicinal property, but rational utilization

should be planned.^[2] *Scoparia dulcis* Linn. locally known as *Manithumbe Gida* belonging to family *Scophularaceae* is a popularly used antiurolithiatic drug in southern parts of India.^[3] It is also known as *Kallurukki* i.e. stone crusher in *Malyalam* language.^[4] It is an erect, usually many branched, annual plant growing 50-100cm tall, tropical plant. This weed grows in moist thickets or along sandy stream beds, commonly gathered from wild.^[5]

It is indigenous to tropical America but commonly found as a weed in Bengal and Tamilnadu and in many parts of India.^[6] Whole plant decoction is used in urinary calculi by traditional practitioners. An antidiabetic compound, amellin is also reported from leaves and stem of the plant, and is used in anemia, albuminuria, ketonuria and other complications associated with diabetes mellitus.^[7] Such an important plant used by traditional community, but not yet recorded quality parameters. Hence an attempt to establish quality parameters of this matured medicinal weed, same presented in this paper.

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MATERIALS AND METHODS

Matured plant of *Scoparia dulcis* Linn. were collected from Udupi District, cleaned properly, authenticated using floras, and sample deposited at SDM Centre for Research in Ayurveda and Allied Sciences Udupi (Voucher No:859/17011801). Plant was washed properly in tap water and shade dried. After complete air drying the plant material was powdered and preserved for further study. Some fresh samples were preserved in FAA solution for microscopic study.

Macroscopy

The external features of the test samples were documented using Canon IXUS digital camera. Organoleptic features of flowers like colour, taste, appearance, smell were recorded according to standard guidelines.^[8]

Microscopy

The preserved specimens were cut into thin transverse section using a sharp blade and the sections were stained with saffranine. The slides were also stained with iodine in potassium iodide for detection of starch. Transverse sections were photographed using Zeiss AXIO trinocular microscope attached with Zeiss AxioCam camera under bright field light. Magnifications of the figures are indicated by the scale-bars.^[9]

Powder microscopy

A pinch of powder was warmed with drops of chloral hydrate on a microscopic slide and mounted in glycerine. Slides were observed, photographed and characteristic features marked.^[9]

Physicochemical standards

The percentage of foreign matter, loss on drying, total ash and acid insoluble ash were determined according to the method described in Indian Pharmacopeia and the WHO guidelines on quality control methods for medicinal plants materials.^[10]

Preliminary phytochemical screening

Preliminary phytochemical screening of the bark powder was performed using alcoholic extract to detect the presence of secondary metabolites.^[11]

HPTLC

1g. of *Scoparia dulcis* Linn. (whole plant) powder was extracted with 10 ml of alcohol. 3, 6 and 9 μ l of the above extract was applied on a pre-coated silica gel F254 on aluminum plates to a band width of 7 mm using Linomat 5 TLC applicator. The plate was developed in Toluene : Ethyl acetate (7.0 : 1.0). The developed plates were visualized in UV 254, 366, and then derivatised with vanillin sulphuric acid and scanned under UV 254nm, 366nm and 620nm. R_f, colour of the spots and densitometric scan were recorded.^[12]

RESULT

Macroscopy

S. dulcis Linn. is a small branched herb found throughout India as a weed. Leaves are serrate 1.3 – 3.8 cm sub acute at apex. The leaves are dark green in colour with pubescence and petiole 9mm in length. Both the surfaces of the leaves having pinnate venation and having prominent veins. Flowers 3-6 from each whorl on slender 8-13 mm long pedicels, white colour with 4-5 sepals. It has a pungent odour. It is bitter in taste.



Fig. 1: Macroscopy of *S. dulcis*

Microscopy

TS of leaf

Transverse section of dorsiventral leaf showed thick upper epidermis, and lower epidermal cells. Between two epidermal layers, mesophyll was differentiated into palisade and spongy parenchyma. Spongy parenchyma had few layered loosely arranged cells. Mesophyll layer showed few rosette calcium oxalate crystals. Bicollateral vascular bundle was found with xylem on the inner side and phloem on the outer side. (Figure 2)

Fig. 2: TS of leaf *S. dulcis* Linn.

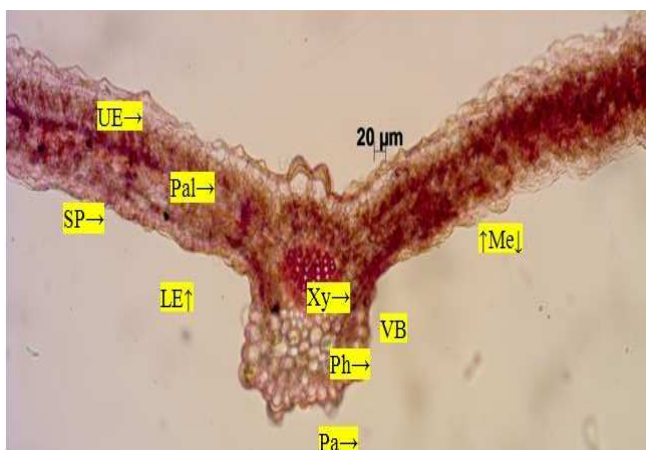


Fig 2a. T.S of *S. dulcis* Linn.

LE - lower epidermis; Me - mesophyll; Pal - palisade; Ph - phloem; SP - spongy parenchyma; UE - upper epidermis; VB - vascular bundle; Xy - xylem.

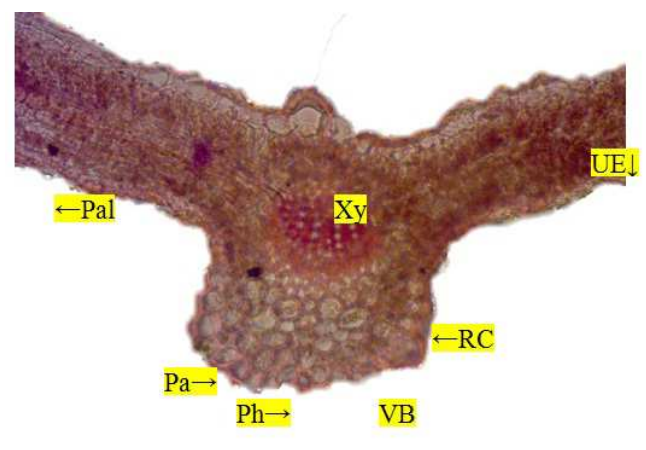


Fig 2b. T.S of leaf enlarged

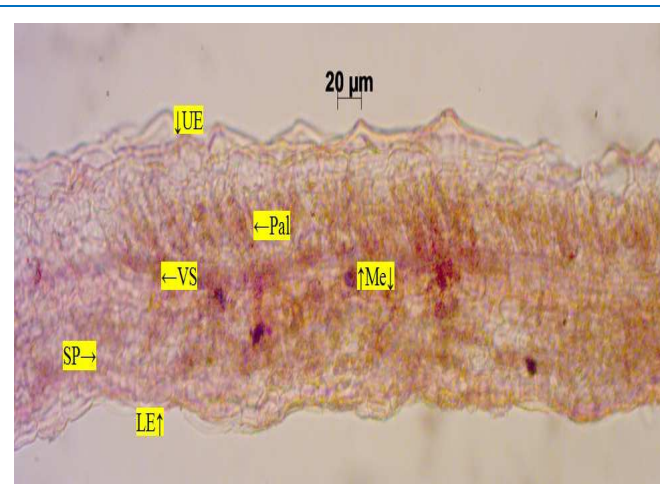


Fig 2c. Lamina enlarged

LE - lower epidermis; Me - mesophyll; Pa - parenchyma; Pal - palisade; Ph - phloem; RC - rosette crystal; SP - spongy parenchyma; UE - upper epidermis; VB - vascular bundle; VS - vascular strand; Xy - xylem.

Fig. 3: Microscopy of matured stem of *S. dulcis* Linn.

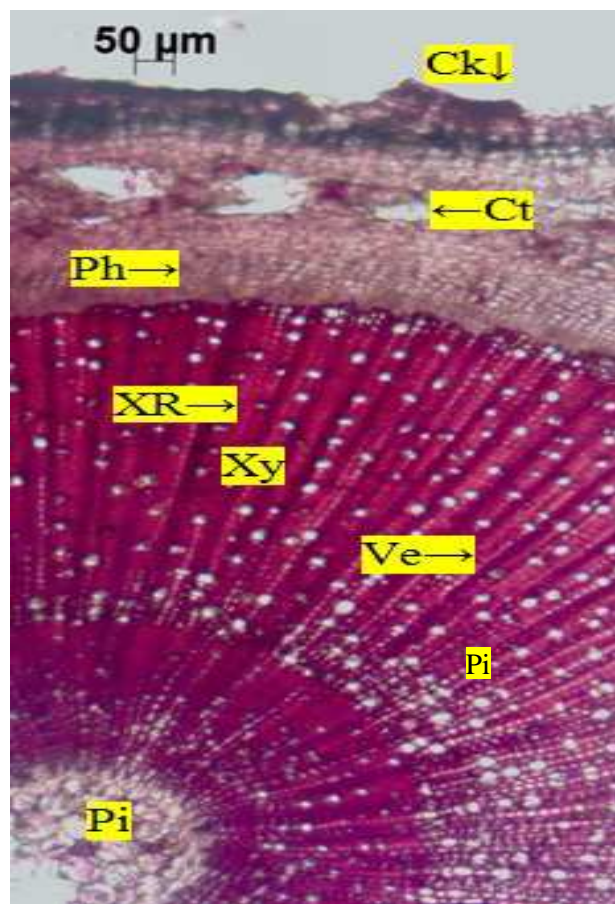


Fig 3a. T.S of stem

TS of stem

TS of stem is almost oval in outline. It shows outer cork tissue, a layer of cortex followed by phloem and rays of xylem surrounding central pith. Outer cork cells are tangentially arranged, few layered. Cells of cortex are parenchymatous and few cholenchymatous, bearing starch grains and stone cells. Few sclerenchymatous cells are also found. Phloem parenchyma surrounds inner xylem rays. Vascular bundles are conjoint collateral closed, with xylem on inner side and phloem on outside. Central pith cells are of parenchymatous with thick cell wall. (Figure 3,4)

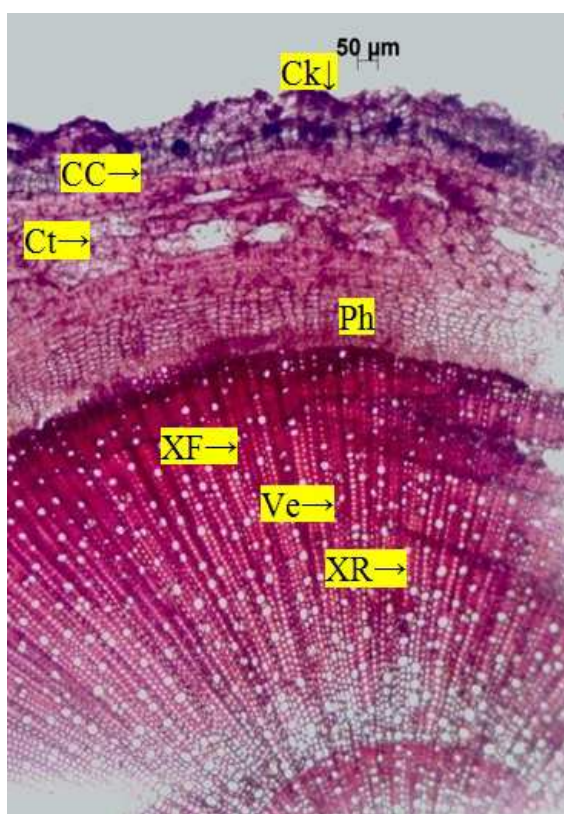


Fig 3b. T.S of stem enlarged
 CC - cork cambium; Ck - cork; Ct - cortex; Pa - parenchyma; PF - phloem fibres; Ph - phloem; Pi - pith; XF - xylem fibres; XR - xylem rays; Ve - vessel.

Powder microscopy

Powder shows fragments of fibres, pitted vessels, reticulate and spiral vessels. Parenchyma cells with and without starch grains are also found. Bunch of starch grains, anther, ovary and pollen grains are marked features of powder microscopy. (Figure 5)

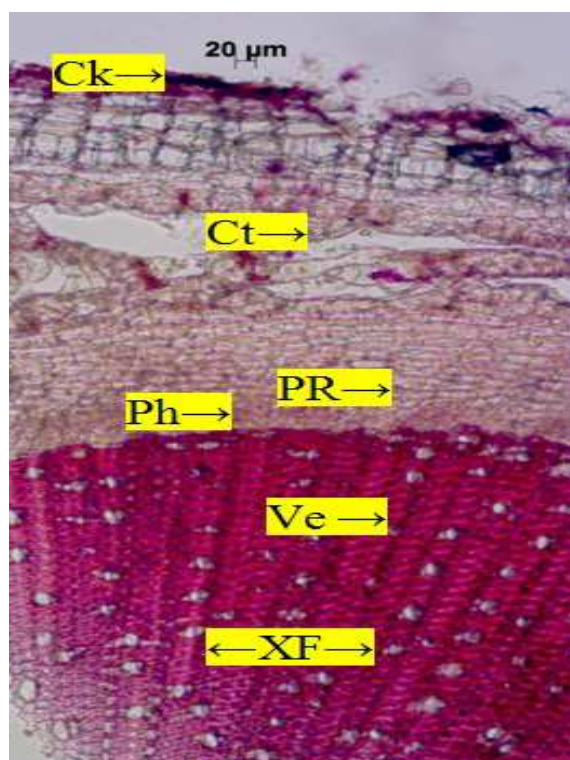


Fig 3c. T.S of stem enlarged

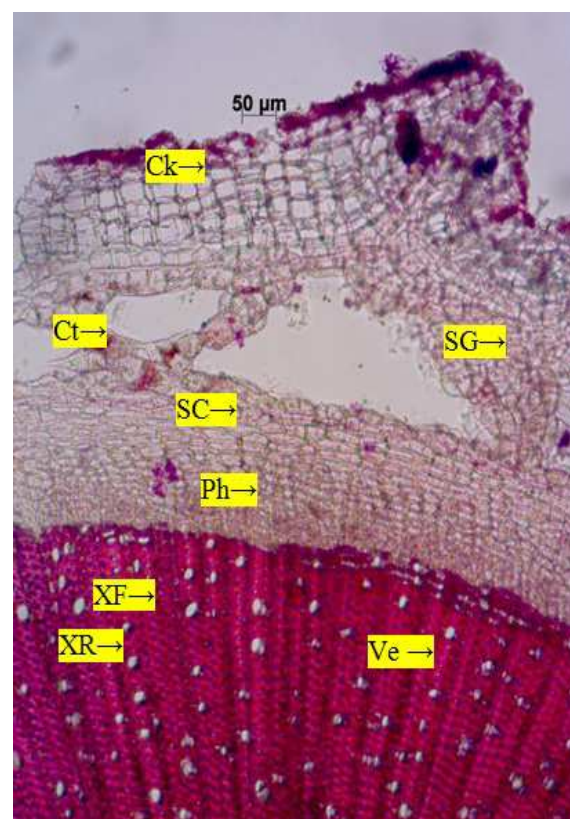


Fig 3d. Cork, Cortex, xylem
 Ck - cork; Ct - cortex; Pa - parenchyma; PF - phloem fibres; Ph - phloem; SG - starch grains; SC - stone cells; Ve - vessel; XF - xylem fibres; XR - xylem rays.

Fig. 4: Microscopy of young stem of *S. dulcis* Linn.

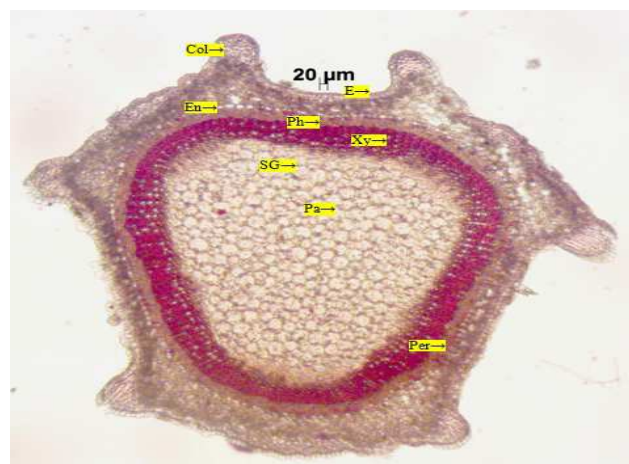
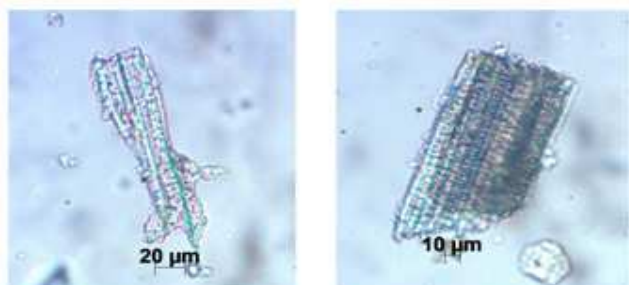


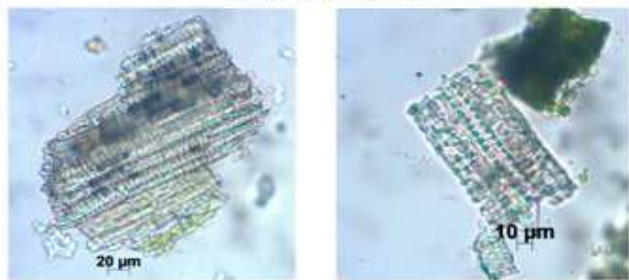
Fig 4a. T.S of Young Stem

Col – Collenchyma; En – endodermis; E – epidermis; Pa – parenchyma; Per – pericyclic fibres; Ph – phloem; SG – starch grains; Xy–xylem.

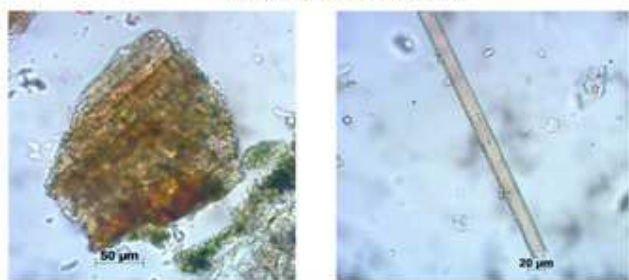
Fig. 5a. Powder microscopy of *Scoparia dulcis* (Whole plant)



Pitted and spiral vessels

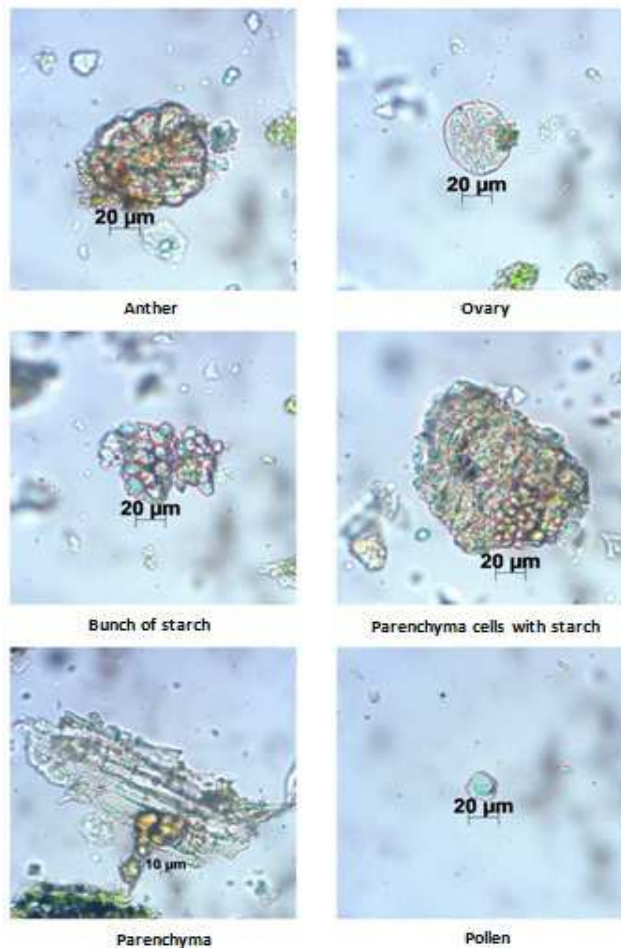


Pitted and reticulate vessels



Vessels

Figure 5b. Powder microscopy of *Scoparia dulcis* (Whole plant)



Physico-chemical standards

S. dulcis Linn. whole plant powder was tested for loss on drying, total ash, acid insoluble ash, ethanol and water soluble extractive as per standard protocol. Loss on drying was 12.44 %w/w, total ash was 7.92% w/w, Acid insoluble ash 0.60%w/w, Water soluble ash 3.29% w/w, alcohol soluble extractive 7.97%w/w, water soluble extractive value 2.47%w/w. Physicochemical standards were a representative of its purity, physical nature and chemical composition. (Table 1)

Table 1: Physicochemical standards of *Scoparia dulcis* Linn.

Parameter	Results n = 3 %w/w
Loss on drying	12.44
Total Ash	7.92

Acid Insoluble Ash	0.60
Water soluble Ash	3.29
Alcohol soluble extractive value	7.97
Water soluble extractive value	2.47

Phytochemical study

Herbs are effective in particular therapeutics because of their chemical nature. Preliminary phytochemical test will decide basic chemical nature of the drug. The test drug powder has shown the presence of alkaloids, carbohydrates, tannin and coumarins.(Table 2)

Table 2: Preliminary phytochemical results of *Scoparia dulcis* Linn.

Test	Inference
Alkaloid	+
Steroid	-
Carbohydrate	+
Tannin	+
Flavanoids	-
Saponins	-
Terpenoid	-
Coumarins	+
Phenols	-
Carboxylic acid	-
Amino acids	-
Resin	-
Quinone	-
(+)- present; (-) – negative	

HPTLC

HPTLC finger print profile of ethanolic extract of *S. dulcis* has been obtained with Toluene: Ethyl acetate (7.0: 1.0) solvent system. The developed plates were

visualized under UV light and white and then under light after derivatisation with vanillin sulphuric acid reagent. R_f, colour of the spots and densitometric scan at 254 and 366nm were recorded. On photodocumentation there were 11 spots under short UV, 15 at long UV, and 9 spots after post derivatisation(Figure 6, Table 3). Densitometric scan at 254 nm showed 13 peaks whereas at 366nm showed 17 peaks (Figure 7)

Fig. 6. HPTLC photo documentation of ethanol extract of *S. dulcis* Linn.

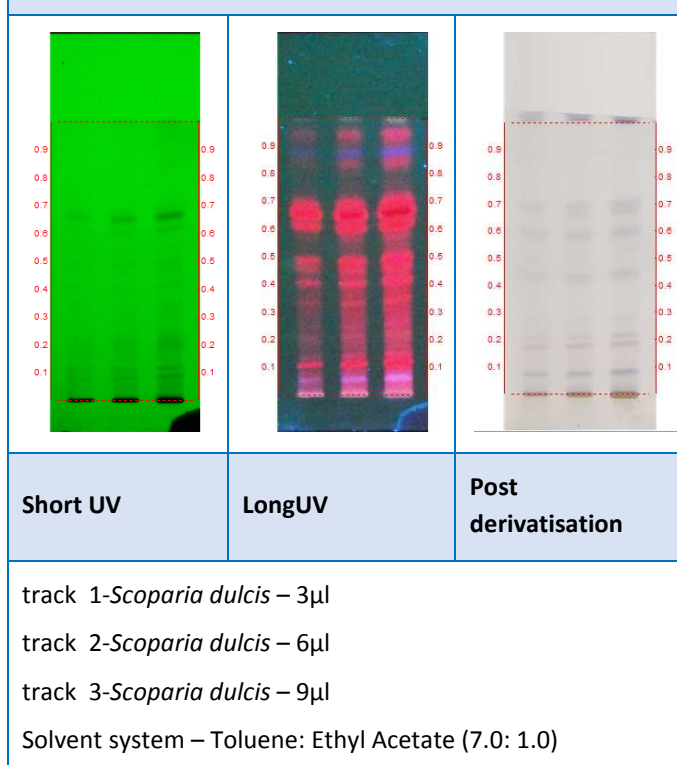
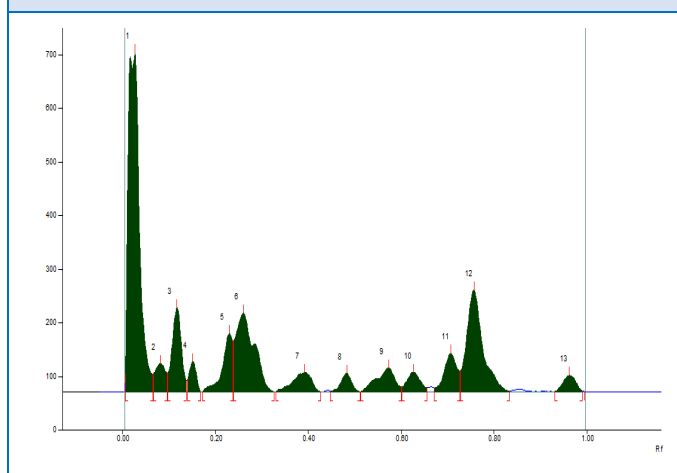


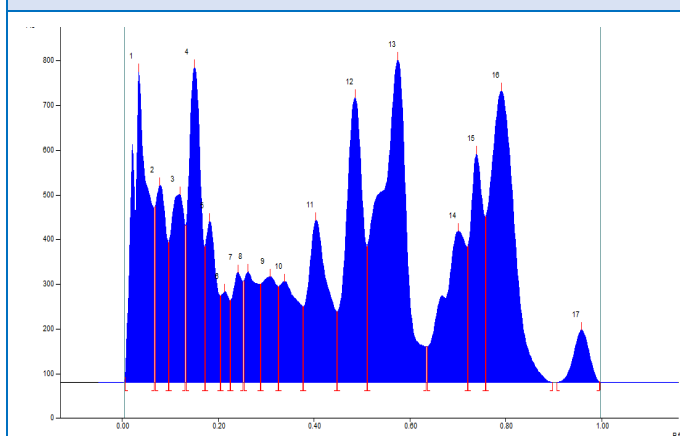
Fig. 7. Densitometric scan of *S. dulcis* Linn.



Track 3. ID: *Scoparia dulcis*

Peak	Start Position	Start Height	Max Position	Max Height	Max %	End Position	End Height	Area	Area %
1	0.01 Rf	33.0 AU	0.03 Rf	632.3 AU	39.48 %	0.07 Rf	33.4 AU	11278.0 AU	35.95 %
2	0.07 Rf	34.1 AU	0.08 Rf	53.0 AU	3.31 %	0.10 Rf	35.4 AU	851.2 AU	2.71 %
3	0.10 Rf	35.5 AU	0.12 Rf	157.1 AU	9.81 %	0.14 Rf	21.3 AU	2271.5 AU	7.24 %
4	0.14 Rf	22.1 AU	0.15 Rf	56.7 AU	3.54 %	0.17 Rf	0.8 AU	628.9 AU	2.00 %
5	0.17 Rf	0.2 AU	0.23 Rf	109.7 AU	6.85 %	0.24 Rf	94.9 AU	1717.2 AU	5.47 %
6	0.24 Rf	95.0 AU	0.26 Rf	146.7 AU	9.16 %	0.33 Rf	0.1 AU	4196.6 AU	13.38 %
7	0.33 Rf	0.3 AU	0.39 Rf	36.7 AU	2.29 %	0.43 Rf	0.1 AU	1074.0 AU	3.42 %
8	0.45 Rf	1.6 AU	0.48 Rf	34.8 AU	2.17 %	0.51 Rf	0.5 AU	575.7 AU	1.84 %
9	0.51 Rf	0.8 AU	0.57 Rf	44.5 AU	2.78 %	0.60 Rf	9.3 AU	1261.0 AU	4.02 %
10	0.60 Rf	9.3 AU	0.63 Rf	37.2 AU	2.32 %	0.66 Rf	8.2 AU	762.9 AU	2.43 %
11	0.67 Rf	7.1 AU	0.71 Rf	72.3 AU	4.51 %	0.73 Rf	38.4 AU	1420.3 AU	4.53 %
12	0.73 Rf	38.8 AU	0.76 Rf	189.9 AU	11.85 %	0.83 Rf	1.4 AU	4714.0 AU	15.03 %
13	0.93 Rf	1.1 AU	0.96 Rf	30.7 AU	1.92 %	0.99 Rf	2.4 AU	621.0 AU	1.98 %

Fig 7a. At 254nm



Track 3. ID: *Scoparia dulcis*

Peak	Start Position	Start Height	Max Position	Max Height	Max %	End Position	End Height	Area	Area %
1	0.01 Rf	10.8 AU	0.04 Rf	696.0 AU	9.78 %	0.07 Rf	90.8 AU	16002.9 AU	9.08 %
2	0.07 Rf	393.7 AU	0.08 Rf	440.4 AU	6.19 %	0.10 Rf	12.7 AU	7083.1 AU	4.02 %
3	0.10 Rf	314.8 AU	0.12 Rf	420.0 AU	5.90 %	0.13 Rf	50.8 AU	8468.9 AU	4.80 %
4	0.13 Rf	351.3 AU	0.15 Rf	704.7 AU	9.90 %	0.17 Rf	03.6 AU	13003.5 AU	7.38 %
5	0.17 Rf	305.4 AU	0.18 Rf	359.9 AU	5.05 %	0.20 Rf	93.0 AU	5724.7 AU	3.25 %
6	0.21 Rf	193.1 AU	0.21 Rf	202.6 AU	2.85 %	0.23 Rf	83.0 AU	2536.5 AU	1.44 %
7	0.23 Rf	183.4 AU	0.24 Rf	245.7 AU	3.45 %	0.25 Rf	25.7 AU	3779.7 AU	2.14 %
8	0.25 Rf	227.5 AU	0.26 Rf	247.3 AU	3.47 %	0.29 Rf	18.9 AU	5061.5 AU	2.87 %
9	0.29 Rf	219.4 AU	0.31 Rf	236.8 AU	3.33 %	0.33 Rf	14.5 AU	5234.0 AU	2.97 %
10	0.33 Rf	214.7 AU	0.34 Rf	225.6 AU	3.17 %	0.38 Rf	69.6 AU	6426.5 AU	3.65 %
11	0.38 Rf	169.8 AU	0.40 Rf	363.3 AU	5.10 %	0.45 Rf	58.3 AU	10874.8 AU	6.17 %
12	0.45 Rf	158.9 AU	0.49 Rf	637.5 AU	8.95 %	0.51 Rf	03.5 AU	16004.2 AU	9.08 %
13	0.51 Rf	305.8 AU	0.58 Rf	722.0 AU	10.14 %	0.64 Rf	79.6 AU	28457.5 AU	16.14 %
14	0.64 Rf	79.7 AU	0.70 Rf	338.1 AU	4.75 %	0.72 Rf	02.3 AU	11778.3 AU	6.68 %
15	0.72 Rf	305.8 AU	0.74 Rf	511.0 AU	7.18 %	0.76 Rf	70.1 AU	9687.9 AU	5.50 %
16	0.76 Rf	371.3 AU	0.79 Rf	651.9 AU	9.16 %	0.90 Rf	0.1 AU	23470.4 AU	13.32 %
17	0.91 Rf	0.2 AU	0.96 Rf	116.9 AU	1.64 %	1.00 Rf	1.0 AU	2672.3 AU	1.52 %

Fig 7b. At 366nm

Table 3: R_f values of samples

Short UV (254nm)	Long UV (366nm)	Post derivatisation
0.06 (L. green)	0.06 (FL. pink)	-
0.08 (L. green)	-	0.08 (L. purple)
0.12 (L. green)	0.12 (FD. red)	-
-	0.14 (FD. red)	-

0.18 (L. green)	-	0.18 (L. purple)
0.21 (L. green)	0.21 (FD. red)	0.21 (L. purple)
-	0.25 (FD. red)	-
-	-	0.28 (L. purple)
-	0.30 (FD. red)	-
0.35 (L. green)	0.35 (FD. red)	0.35 (L. purple)
-	0.40 (FD. red)	-
0.42 (L. green)	-	0.42 (L. purple)
-	0.44 (FD. red)	-
0.48 (L. green)	0.48 (FD. red)	-
0.50 (L. green)	-	-
-	0.60 (FD. red)	0.60 (L. purple)
0.62 (L. green)	-	-
0.67 (D. green)	0.67 (FD. red)	0.67 (L. purple)
-	-	0.70 (L. purple)
-	0.85 (FD. red)	-
-	0.88 (FD. violet)	-
-	0.95 (FD. red)	-

F - Fluorescent; L - Light; D - Dark

DISCUSSION

Scoparia dulcis Linn. a member of *Scophularaceae* family is a less known drug, but used by traditional practitioners in the treatment of urinary calculi. This is a perennial herb with woody stem commonly called as *Manitumbegida*. Scientifically recorded authentic standard parameters for herbs are an essential step in its global promotion as well as in clinical competence.^[13] *S.dulci* Linn. provided with macro-microscopic illustrations, physicochemical standards, an inference of secondary metabolites and HPTLC are indicative of its quality standards.

TS of leaf has shown the presence of mesophyll which is differentiated into palisade and parenchymal layer, in between two epidermal layers. Bicollateral vascular bundles and presence of rosette calcium oxalate crystals at mesophyll layer are characteristic features of Leaf anatomy. Outer cork tissue, a layer of cortex

consisting of parenchymatous and cholenchymatous cells possessing starch grains, stone cells are features of stem histology. Conjoint collateral closed vascular bundles and central pith are inclusions of stem TS. Powder shows fragments of fibres, pitted vessels, reticulate and spiral vessels, starch grains, anther, ovary and pollen grains. Physico-chemical standards are a measure of purity, and physical nature.^[14] High loss on drying value (12.44 %w/w) indicates moisture content, whereas total ash value (7.92% w/w) shows total carbonaceous matter. Alcohol soluble extractive was much better (7.97%w/w) than water soluble extractive value (2.47%w/w). Alkaloids, carbohydrates, tannin and coumarins were main secondary metabolites detected out of test drug. HPTLC fingerprints are a record of its different chemical constituents. Thus the quality monograph prepared on this drug beneficial in future research.

CONCLUSION

Growing herbal drug industry necessitates quality parameters for raw material for its global acceptance. *S. dulci* Linn. with scientifically recorded macro-microscopic features, along with other quality parameters forms authentic identity of the drug, thus prevent any admixture.

REFERENCES

- Dintu Jose, Pharmacognostical and Experimental study on Arikal Tumbe- *Anisochilus carnosus* (L.f.) Wall. wsr to its Antifungal activity, MD Thesis, RGUHS Bangalore;2017.
- Mallya Suma V, Comprehensive study of Plants in Surasaadi Gana wsr o their antibacterial activity (PhD thesis), Nashik, Maharashtra: Maharashtra University of Health Sciences; 2016.
- Anonymous 1972-1999, Wealth Of India, Council for Scientific And Industrial Research vol-9, Published by Publication and Information Directorate, New Delhi 1972, Pp 472, Page no: 260- 261
- Kirtikar KR & Basu BD.Indian Mediicnal Plants, Volume III, Dehradun; International Book Distributors; 2008 p 2320
- Bhat Gopalakrishna K., Flora Of Udupi, Published by Indian Naturalist, Udupi, Edition 2003, Pp 913, Page no: 422
- Hooker J. D., Flora Of British India, Vol- 4, Published under The Authority Of The Secretary Of State For India in Council, Pp 780, Page no: 289
- Khare CP. Indian Medicinal Plants, An illustrated dictionary. New Delhi; Springer;2007;p 592
- Juliet et al. Authentication and Proximate analysis of *Madhuka indica* Gmell. A wild edible flowers used for traditional fermentation of Ayurvedic Biomedicines; J Ayu Med Sci 2016;1(1);12-18
- Dintu et al. Pharmacognostical evaluation of an extrapharmacopeial Drug of Ayurveda- *Anisochilus carnosus*(L.F.)Wall. J Ayu Med Sci 2017;2(2): p179-87
- Mallya Suma V, Nesari Tanuja, Sunilkumar KN, Pharmacognostic standards of Katphala(*Myrica nagi* Hook.f.non-Thumb); A potent bark drug used in Indian system of medicine. Journal of Scientific and innovative Research 2016;5(4): p1-3
- Quality control methods for medicinal plants. Geneva: World Health Organization; 1998; p25-8
- Brain KR, Turner T, The practical evaluation of phytopharmaceuticals , Bristol; Wright-Scientechnia;1975; p. 10-12
- Sethi PD, High Performance Thin Layer Chromatography, 1st edition, New delhi; CBS Publishers and Distributors; 1996; p1-56.
- Mukharjee Pulok K. Quality Control of Herbal Drugs. New Delhi: Business Horizons;2002;p68
- Joshi Devendra, Joshi Geeta. Quality Control &Standardization of Ayurvedic Medicines. Varanasi: Chaukamba Orientalia; 2011; p113

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