IJPSR (2023), Volume 14, Issue 12



INTERNATIONAL JOURNAL

(Research Article)



Received on 15 April 2023; received in revised form, 16 June 2023; accepted 04 July 2023; published 01 December 2023

PHARMACOGNOSTIC AND HPTLC COMPARISON OF KERALA MARKET SAMPLES OF BHARANGI (CHERUTHEK) WITH GENUINE SOURCE PLANT *ROTHECA SERRATA* (L.) STEANE & MABB

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Keywords:

Bhārangī, (*Rotheca serrata* (*L*.) Steane & Mabb, Pharmacognosy, Market study, Adulteration, *Ayurveda*

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ABSTRACT: Background: Bhārangī, herbal drug used in Ayurveda as Kapha-Vata samaka and indicated in Swasa, Kasa, Vrana, Shotha etc. It is an important ingredient in many of the preparations like Kantakaryavaleha, Kanakasava, Dasamularista, Lodhrasavam, Sudarsanasavam. As per API the accepted source of Bhārangī is Rotheca serrata (L.) Steane & Mabb belongs to family Lamiaceae. There were ambient studies in which it was found that roots of Bhārangī had been adulterated with stem of the same in the market. But there were no published works indicating the presence of adulterants in the Kerala market, hence this study was proposed to assess the genuineness of raw drug available in the name of Cheruthek $(Bh\bar{a}rang\bar{i})$ in different Kerala markets with the root of genuine source plant R. serrata. Materials and Methods: Fresh root of Bhārangī (Rotheca serrata (L.) Steane & Mabb) are collected from the natural habitat. The market samples of Cheruthek were collected from six different regions in Kerala. All samples were screened for pharmacognostic parameters- macroscopic, microscopic, histo-chemical and powder microscopic characterization along with HPTLC profile and compared with the genuine drug. Result & Conclusion: After the pharmacognostic comparison and HPTLC profiling of six market samples of the drug Cheruthek collected from selected districts of Kerala, the Kozhikode sample had similar characters of the R. serrata; Thiruvananthapuram sample was having characters of the plant Premna herbaceae Roxb. All other samples were adulterated with various plant parts.

INTRODUCTION: *Ayurveda* is increasingly gaining recognition as one of the medical sciences worldwide in the last few decades. Plant materials were used throughout the world as home remedies, over-the-counter drug products and raw materials for the pharmaceutical industry, and represent a substantial proportion of the global drug market. Commercialization in the medical system leads to substituting or adulterating herbal drugs to withstand scarcity.

QUICK RESPONSE CODE	DOI: 10.13040/IJPSR.0975-8232.14(12).5676-86	
	This article can be accessed online on www.ijpsr.com	
DOI link: https://doi.org/10.13040/IJPSR.0975-8232.14(12).5676-86		

Using commercially available inferior-quality drugs had compromised the therapeutic efficacy of *Ayurveda* medicine. Hence, the quality standardization of drugs had got greater importance in *Ayurveda*. As per CCRAS, a market sample survey has been found to be one of the priority areas for drug research 1 .

Bhārangī, a potent Ayurvedic drug used in respiratory disorders, which acts as Kapha Vata Samaka. In classics the roots were considered as the part used along with root bark which is used in many of the Ayurvedic formulations like Bharangyadi kashaya, Bharangyadi churna, Kantakaryavaleha, Bhārangī Guda, Kanakasava, Lodhrasava etc. As per Ayurvedic Pharmacopeia of India and Quality Standards of Indian Medicinal

Plants, states that Rotheca serrata (L.) Steane & Mabb belongs to the family Lamiaceae is the botanical source of *Bhārangī* and the official part used is dried roots $^{2, 3}$. Due to increased demand and over-exploitation, there is a significant reduction in the availability of genuine drug; hence chances for adulteration with other spurious, inferior or useless parts of same or different plants is very high. Some previous studies found that the roots of *Bhārangī* are adulterated with the stem, in many markets. From a previous study conducted on the market samples of Bhārangī, it was found that all the samples were adulterated and none of the samples were genuine (Singh et. al. 2015)⁴. The most of the samples obtained in South Indian are observed to be the roots and root nodules of Pygmaeopremna herbacea (syn. Premna herbacea) (Vasudevan et. al. 1971)⁵. Though Brihattrayis and Nighantukara have described only one variety of Bhārangī, we come across two types of Bhārangī in Vaidyaka Sabda Sindhu, the white flowered and blue flowered. This description has become a source for controversy. The latter authors clarified that the white-flowered variety is identified as Clerodendrum indicum (L.) Kuntze and the blue flowered one as Rotheca serrata (L.) Steane & Mabb^{6,7}. There were no published works are available about the genuinity of Bhārangī in Kerala market samples. Hence in this study, mainly aimed to compare the different market samples of Bhārangī (Cheruthek) by Pharmacognostic and High-performance thin layer chromatography (HPTLC) profiling with the root of genuine source plant Rotheca serrata (L.) Steane & Mabb.

MATERIALS AND METHODS: The root and stem of genuine sample of Bhārangī (Cheruthek) collected from Herbal Garden were of Vaidyaratnam P. S. Varrier Ayurveda College, Kottakkal and Vrindavanodyanam, Thanal Mathrusadanam, Mayanoor, Thrissur. The samples were authenticated and specimens were deposited at the Herbarium of Center for Medicinal Plants Research (CMPR), Arya Vaidya Sala, Kottakkal, Kerala. (Herbarium Voucher No: 10047). The different market sample of Bhārangī (Cheruthek) were procured from six different markets of Kerala. (Thiruvananthapuram, Idukki. Palakkad. Kozhikode, Wayanad and Kasargod) The genuine sample of Rotheca serrata (L.) Steane & Mabbwere compared with the market samples, on basis of pharmacognostic parameters along with HPTLC profile.

Samples were named as follows:

Samples 1: Roots and stem of *Rotheca serrata* (*L*.) Steane & Mabb

Sample 2: Sample collected from Thiruvananthapuram market.

Sample 3: Sample collected from Idukki market.

Sample 4: Sample collected from Palakkad market.

Sample 5: Sample collected from Kozhikode market.

Sample 6: Sample collected from Wayanad market.

Sample 7: Sample collected from Kasargod market.

Sample 8: Root of *Clerodendrum indicum*.

Methodology: External characters were studied and documented per standard API protocol. The materials for anatomical study were processed as per the standard protocol. The powder analysis and maceration study were done. Observations were done under Leica DM 1000 LED microscope and photographs were taken using Leica DFC 295 inbuilt camera. For HPTLC profiling of the water extracts of the samples were done under the solvent system - Ethyl acetate: methanol: water (9:1:1). Observed under UV light at 254, 366nm and after derivatization with ANS reagent, recorded the R_f value. Densitometric scanning of plates was done using Camag TLC scanner 3 at 254, 366 and 550nm.

RESULTS: The genuine sample of *Bhāraṅgī* and its six market samples were compared by its pharmacognostic parameters using macroscopic, microscopic, histochemical, powder microscopic characters along with HPTLC profile.

Macroscopic Comparison: Organoleptic characters of the Samples of $Bh\bar{a}rang\bar{a}$ were shown in **Table 1** and **Fig. 1**. Samples showed almost identical characteristics such as colour, fractures, external markings and cut surfaces. The main differences observed between the samples were in

thickness. The sample 1; root and stem had extremely different texture compared to market samples. Sample 6 showed extremely different macroscopy when compared to others.

		Colour	Shape	Size	Surface	Fractures	Odour	Taste
Sample	<i>R</i> .	Outer surface	Cylindrical	Diameter	Rough, longitudinal	Short	Characteristic	Acrid
1	serrata–	brown in colour,		upto 2.5cm	striations &		odour	
	Root	wood portion		1	lenticels			
	<i>R</i> .	creamish white	Cylindrical	Diameter	Rough, longitudinal	short	No	Acrid
	serrata–			upto 3cm	striations &		Characteristic	
	Stem			1	lenticels		odour	
Sample	Root	Dark to light	Cylindrical long	Diameter	Rough; longitudinal	Short,	Characteristic	Acrid
2		muddy brown	slender;	0.5-1cm	striations	brittle	odour	
		colour, Wood	stoloniferous					
		portion is	roots					
	Stolon	creamy brown	Cylindrical long	Diameter 2-	Rough; longitudinal	Short,	Characteristic	Acrid
			slender;	3cm	striations	brittle	odour	
			stoloniferous					
			roots					
	Tuber		Cylindrical long	Diameter 2-	Rough; longitudinal	Short,	Characteristic	Acrid
			slender;	3cm	striations	brittle	odour	
			stoloniferous					
			roots					
Sample	Root	Dark to light	Cylindrical,	root pieces-	Rough; longitudinal	Short,	Characteristic	Acrid
3		muddy brown	lateral roots-	1-1.5cm in	striations	brittle	odour	
		colour, Wood	long slender;	diameter; 6-				
		portion is		8cm long;				
	Tuber	creamy brown	stoloniferous	Root	Rough; longitudinal	Short,	Characteristic	Acrid
			roots,	nodules-	striations	brittle	odour	
			subglobous	Diameter				
a 1				2.5-3cm;				
Sample	Root	Brown in	Cylindrical,Late	0.5-1cm in	Rough; longitudinal	Short	Characteristic	Acrid
4		colour; Wood	ral roots were	diameter, 3-	striations;minute		odour	
		portion is	long slender;	4cm long	lenticels			
	Charma	creamy brown		pieces	Develo levelo divel	Ch aut	Channataniatia	له ند ۸
	Stem	Brown in colour	stout to slender,	diameter of 2-2.5 cm	Rough; longitudinal striations; minute	Short	Characteristic odour	Acrid
			cylindrical	2-2.5 CIII	lenticels		ouour	
Commlo	Root	Brown in	Cylindrical,	1-1.5cm in		Short	Characteristic	Acrid
Sample 5	KUUL	colour; Wood	Lateral roots	diameter, 5-	Rough, longitudinal striations	Short	odour	Achu
5		portion is	were long	7cm long	sulations		ououi	
		creamy brown	slender	pieces				
	Stem	Brown in colour	Stout to slender,	Diameter of	smooth;longitudinal	Short	Characteristic	Acrid
	Stem	Diowii in coloui	cylindrical	2-2.5cm	striations, minute	Short	odour	Actiu
			cymarical	2 2.5em	lenticels		odour	
San	ple 6	Dark- light	Fibrous	1.5-2cm	Rough -smooth		No	
Suit	-p-• •	brown in colour,	11010005	long pieces	itougii sinootii		Characteristic	
		,		8 F			odour	
Sample	Root	Brown in	cylindrical;	0.5-1cm in	Rough longitudinal	Short	Characteristic	Acrid
7		colour, wood	Lateral roots	diameter,1-	striations		odour	
		creamy brown	were long	2cm long				
			slender;	pieces				
	Stem	Brown in colour	stout to slender,	Diameter of	Smooth,	Short	Characteristic	Acrid
			cylindrical&	1-2cm	longitudinal		odour	
			branched		striations			
C. indic	um - Root	Yellowish	Cylindrical,	1cm in	Smooth;	Short	Characteristic	Acrid
		brown in colour;		diameter	longitudinal		odour	
		wood portion			striations			
		creamish white						

TABLE 1: MACROSCOPIC COMPARISON OF THE SAMPLES OF BHĀRAṅGĪ

Microscopic Comparison:

Microscopy of Sample 1 (Root and Stem):

Root: TS of the root was almost circular in outline with outer layers of cork ruptured at many places followed by a wide cortex, phloem and welldeveloped xylem region. Detailed TS showed a well-developed cork region with two distinct zones - outer zone consisting of up to 15 layers of comparatively thick walled rectangular to tangentially elongated cells which often peeled off at many places. Inner zone composed of more than 15 layers of tangentially elongated radially arranged cells. Cortex was composed of 10 - 15 layers of oval or elongated loosely arranged parenchymatous cells with inter-cellular spaces. Simple and compound starch grains measuring upto 10-15µm in diameter are scattered throughout the cortical region along with acicular and rod-shaped calcium oxalate crystals.

Solitary and groups of fibers and stone cells present throughout the cortical region. Phloem is comparatively narrow composed of compactly arranged compressed cells often alternating with medullary ray cells at many places. Simple starch grains and acicular and rod-shaped crystals of calcium oxalate were present at some places.

Centre portion occupied by xylem elements such as vessels, fibers, parenchyma and medullary rays. Xylem vessels were round to oval in shape, 30-100µm diameter usually in solitary and rarely grouped. Medullary rays 2-3 seriated composed of radially elongated pitted parenchymatous cells often filled with starch grains and acicular crystals of calcium oxalate. Starch grains also observed in xylem elements including parenchyma cells and fibers. The presence of growth rings in mature roots was observed in **Table 2** and **Fig. 2**.

Stem: TS of the stem was almost circular in outline with narrow cork layers which were ruptured at many places followed by wide cortex, narrow phloem and well-developed xylem region. Detailed TS showed narrow cork region with 3-5 layers of rectangular to tangentially elongated cells. Cortex were composed of 10-15 layers of oval to elongated loosely arranged parenchymatous cells with intercellular spaces. Pericyclic fibers seen in cortical region in discontinuous patches alternating with solitary or group of stone cells. Starch grains of diameter 10-15µm and acicular crystals present in some cortical cells. Phloem narrow composed of compactly arranged compressed cells often alternating with medullary ray cells. Xylem region composed of vessels, fibers and medullary rays. Xylem vessels round to oval in shape, 40-150µm diameter usually solitary and rarely in grouped and scattered al-throughout the xylem region.

Tyloses present. Xylem fibers thick walled and possess major part of xylem region. Medullary ray uni-bi seriated composed of radially elongated pitted parenchyma cells. Pith is comparatively large and occupies the central region, composed of parenchymatous cells filled with starch grains and acicular and rod-shaped crystals of calcium oxalate at some regions **Table 2** and **Fig. 2**.

Microscopic Comparison with Market Samples: The TS of the Thiruvananthapuram sample and Idukki sample had dissimilar microscopic characteristics with that of genuine sample. The presence of orange-brown deposition at the inner layers of cork cells were evidently seen in the Thiruvananthapuram and Idukki samples.

The presence of stone cells was very few in the cortical region in the Thiruvananthapuram sample and Idukki sample. While its abundant in case of genuine sample. The presence of cortical fibers found were in the root samples of Thiruvananthapuram sample and Idukki sample. Starch grains and acicular crystals were present comparatively less with genuine sample. Presence of oil globules in case of the Thiruvananthapuram sample and Idukki sample. Palakkad and Kozhikode samples had similar microscopic characteristics to genuine samples.

The presence of stone cells and fibers was even in both samples, starch grains and acicular crystals of calcium oxalate. Wayanad and Kasargod samples did not resemble any of the microscopic characters of the genuine sample. The Wayanad sample prominently showed the vascular region which is dissimilar with that of genuine. The Kasargod sample showed different TS of root and stem pieces. It possesses uni-bi seriated medullary rays. Presence oil globules were seen in both samples **Table 2, Fig. 3.**

		Cork	Cortex	Phloem	Xylem
Sample	Rotheca	Two distinct zones-	Wide; round to oval,	Narrow zone;	Round to oval vessels;
1	serrata –	tangentially elongated	loosely arranged	compactly arranged	Multiseriate medullary rays,
	Root	 peeled off at places. 	parenchyma; Rounded	cells; Rounded starch	rarely uni to bi seriate; Starch
		radially arranged;	starch grains; acicular	grains; acicular and	grains in medullary rays and
		tannin depositions	and rod-shaped	rod shaped crystals;	xylem fibres; acicular and rod
		crystals; scattered	parenchyma	shaped crystals in medullary	
			fibres; stone cells and	alternating with ray	rays. Bigger sized vessels in
			sclereids; tannin	cells	outer and centre portion of the
			deposition		wood
	Rotheca	Narrow; tangentially	Wide; oval to	Narrow zone;	Major portion occupied by
	serrata –	elongated cells; tannin	elongated loosely	compactly arranged	fibres; round to oval vessels; uni
	Stem	depositions	arranged	cells; parenchyma	to bi seriate medullary rays;
			parenchymatous cells;	alternating with ray	Widepith polygonal
			discontinuous ring of	cells	parenchymatous with starch
			pericyclic fibres; stone		grains, acicular and rod shaped
			cells; starch grains and		crystals
			acicular crystals; tannin		
			depositions		
Sample	Root	2-5 layers of	10-12 layers; loosely	Narrow and	round shaped vessels; Bigger
2		tangentially elongated	arranged	compressed; 3-5	sized vessels located at outer
		irregular cells;	parenchymatous cells;	layers; phloem fibers	portion of wood, uni-bi seriate
			cortical fiber patches;		medullary rays;
			prismatic crystals;		
	Stolon	tangentially elongated	5 - 8 layers of	5 - 10 layers of	In radial patches; vessel
		cells; inner layer of	elongated parenchyma	closely packed cells	diameter; alternating with
		cells contains orange	cells; stone cells;		multiseriate medullary rays;
		brown depositions;	tannin depositions		polygonal parenchymatous pith
	Tuber	20 - 25 layers of thick	oval to round	round to oval	multi seriate medullary rays;
		walled, tangentially	tangentially elongated	tangentially	acicular crystals and starch
		elongated cells; inner	thin-walled	elongated cells;	grains; rounded parenchymatous
		layer of cells contains	parenchymatous cells;	parenchyma	pith with starch grains
		orange brown	stone cells; sclereids;	alternating with ray	
		depositions	acicular crystals; tannin	cells; acicular crystals	
			depositions		
Sample	Root	10 - 15 layers, narrow	oval to rectangular	narrow and	Vessels less in number multi-
3		compressed cells; inner	elongated	compressed round to	seriated medullary rays; starch
		layers contain orange	parenchymatous cells;	oval cells;	grains
		brown depositions; oil	Stones cells and	parenchyma	
		globules	sclereids prismatic and	alternating with ray	
			acicular crystals of	cells; acicular crystals	
			calcium oxalate;	of calcium oxalate,	
				presence of oil	
	T 1	4	1 (1	globules	X7
	Tuber	tangentially elongated	oval to round	Narrow and	Vessels less in number; uni - bi
		cells; inner layer of	tangentially elongated	compressed,	seriated medullary rays; pith
		cells contains orange	cells; stone cells;	compactly arranged	loosely arranged
		brown depositions;	sclereids; starch grains,	cells	parenchymatous cells, starch
Some1-	Post	presence of oil globules	oil globules	no	grains present in xylem and pith
Sample 4	Root	rectangular to	Narrow and	narrow and	Vessels round to oval in shape;
		tangentially elongated	compressed layers of	compressed,	Tyloses present; multi - seriated
		cells	oval or elongated cells;	compactly arranged	medullary rays
			Acicular, prismatic and rod-shaped crystals;	cells	
			fibers, stone cells scattered		
	Stem	rectangular to	oval to elongated	Narrow; compactly	Xylem vessels large, round to
	Stell	tangentially elongated	loosely arranged cells;	arranged compressed	oval in shape; multi seriated
		cells; Tannin	Fibres, sclereids; stone	cells alternating with	medullary rays; pith is large
		cons, ramm		cons anomating with	mouthary rays, plui is large

TABLE 2: MICROSCOPIC COMPARISON OF THE SAMPLES OF BHĀRANGĪ

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		depositions	cells; Granular masses and acicular crystals	medullary ray cells	
SampleRoottwo distinct zones;20-30 layers;5outer zones- 8-10elongated lo1ayers; rectangular toarrangedtangentially elongatedparenchymatorcells; - peeled off atSimple and corplaces; Inner zone -starch grains; amore than 5 - 8 layers;and rod-shatangentially elongatedcrystals; store ocells.; tanninfibers; tan		20-30 layers; oval or elongated loosely arranged parenchymatous cells; Simple and compound starch grains; acicular and rod-shaped crystals; stone cells and fibers; tannin depositions	Narrow; compactly arranged compressed cells alternating with medullary ray cells; starch grains; acicular and rod-shaped crystals	Xylem vessels were round to oval in shape, Tyloses present; multi- seriate medullary rays; starch grains and acicular crystals;	
	Stem	Narrow; 8 - 10 layers rectangular to tangentially elongated cells; tannin depositions	15 - 20 layers of oval to elongated loosely arranged parenchymatous cells; pericyclic fibers, stone cells; Granular masses and acicular crystals; tannin depositions	Narrow; compactly arranged compressed cells; cells alternating with medullary ray cells.	vessels were round to oval in shape; Tyloses present; major portion occupied by xylem fibers; uni - bi seriated medullary rays; acicular crystals; pith large; starch grains, prismatic, rod and acicular crystals
San	nple 6	Two distinct zones; reddish brown depositions	Narrow, elongated irregular cells	Wide, compressed ells; with brown depositions; alternating with ray cells; fibres scattered throughout	Well-developed xylem resembling a heart wood. Vessel's solitary with diameter. Medullary raysuni– multi seriate
Sample 7	Root	rectangular to tangentially elongated cells; peeled off at many places	oval or elongated loosely arranged parenchymatous cells; acicular and rod- shaped crystals	Narrow; compactly arranged compressed cells; fibres in solitary and in groups; acicular crystals	vessels were round to oval in shape; uni-bi seriated medullary rays; starch grains
	Stem	Narrow; rectangular to tangentially elongated cells;	oval to elongated loosely arranged parenchymatous cells; Pericyclic fibers in patches; stone cells; Starch grains & acicular crystals	Narrow; compactly arranged compressed cells	vessels were round to oval; uni- bi seriated medullary rays; acicular crystals; starch grains
Sample 8	C. indicum - Root	Two distinct zones; outer zone - upto 5 layers- thick walled rectangular to tangentially elongated cells; peeled off at many places; Inner zone- 5 layers; tangentially elongated cells; tannin depositions	10-12 layers; oval or elongated loosely arranged parenchymatous cells; prismatic crystals; stone cells; starch grains;	Wide, compactly arranged compressed cells, cells alternating with medullary ray cells; starch grains; prismatic crystals;	vessels were round to oval; uni- bi seriated medullary rays; starch grains

HPTLC Profile: HPTLC chromatograms and densitometric profiles of 8 samples at 254nm, 366nm and 550 nm were recorded. **Fig. 4** the Rf values of separated compounds of each sample were noted. **Table 3** At 254 nm, sample 5 has maximum number of peaks (10 peaks), sample 8 with 8 peaks, and sample 4 with 7 peaks. Rf values of many peaks of these samples were similar

indicating same chemical constituents. *Rotheca* serrata (L.) Steane & Mabb. root sample (sample 1R) had only seven peaks and its stem (sample 1S) with 8 peaks. Sample 6 shows very less peaks. Peak intensities were varied in case sample 2 and sample 3. Sample 5 showed some unique peaks that were observed in the genuine stem and root.

 TABLE 3: Rf VALUE OF SAMPLES

2 07 6 7 7 2 9 8 6 2 2 4 7 3 2 5 5 2 3 4 3 0.0 0. 0.1 0.1 0.0 0.4 0.1 0.0 0.0 0.1 0.1 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0	UV 550 3 4 0.0 0.0 7 4 0.1 0.0 4 8	0nm 5 0.0 3 0.0	6 0.2 2	7 0.0 3	8 0.
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\begin{array}{ccc} 0.0 & 0.0 \\ 7 & 4 \\ 0.1 & 0.0 \end{array}$	0.0	0.2 2	0.0	
2 07 6 7 7 2 9 8 6 2 2 4 7 3 2 5 5 2 3 4 3 0.0 0. 0.1 0.1 0.0 0.4 0.1 0.0 0.0 0.1 0.1 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0	7 4 0.1 0.0	3	2		0.
0.0 0. 0.1 0.1 0.1 0.0 0.4 0.1 0.0 0.0 0.0 0.1 0.1 0.0 0.0 0.2 0.0 0.0 0.0 0.0 0.0	0.1 0.0			2	
		0.0		3	04
	4 8		0.2	0.0	0.
7 16 8 5 4 7 4 7 8 7 7 9 6 7 7 8 8 6 9 8 7		7	5	8	08
0.1 0. 0.2 0.1 0.1 0.1 0.2 0.1 0.1 0.1 0.3 0.1 0.1 0.1 0.5 0.1 0.0 0.1 0.1 0.1	0.1 0.1	0.1	0.2	0.1	0.
9 19 7 8 8 5 8 5 9 9 0 8 5 5 9 8 8 5 4 4	9 1	4	9	2	14
0.3 0. 0.4 0.2 0.2 0.1 0.4 0.2 0.3 0.2 0.3 0.3 0.1 0.1 0.9 0.2 0.1 0.2 0.1 0.1	0.2 0.1	0.1	0.3	0.1	0.
1 27 9 6 5 8 8 0 1 7 3 0 8 8 3 9 4 0 9 9	7 4	8	5	8	23
0.4 0. 0.5 0.3 0.2 0.2 0.2 0.4 0.3 0.4 0.4 0.2 0.2 0.9 0.2 0.3 0.3 0.2	0.3 0.1	0.2	0.4	0.2	0.
0 31 6 0 9 5 7 1 1 9 0 5 5 3 6 2 1 8	1 9	6	1	3	25
	0.4 0.3	0.3	0.4	0.2	0.
9 41 1 7 0 9 9 1 4 0 9 0 1 2 2 0	1 0	0	4	9	33
	0.4 0.4	0.4	0.4	0.3	0.
4 50 2 9 1 4 0 3 8 9 6 5 0 1	5 0	0	9	1	45
	0.5 0.4	0.4	0.7	0.3	0.
55 7 7 4 2 7 9 5 4	5 7	5	8	5	57
	0.8 0.5	0.5	0.8	0.7	0.
3 5 4 2 2 0 9	1 2	3	1	9	82
	0.9 0.7	0.7	0.9	0.9	0.
3 3 7 2 1	4 8	8	3	3	94
0.8 0.8	0.8	0.8			
4 7	0	0			
0.9 0.9	0.9	0.9			
5 4	3	3			

DISCUSSION: Quality and Safety are fundamental principles in the provision of herbal medicines for health care. Scientific studies estimate herbal product adulteration as 40-60% in India, the most of Indian herbal medicinal products are essentially mixed with one or a few other herbs that could lessen the therapeutic activity of the main ingredients ⁸. Only a few studies are available on quality standards of some market samples.

A study conducted in South Indian Market samples of $Bh\bar{a}rang\bar{a}$, it was noted that the roots of C. serratum was the accepted species for the drug. As per previous study other species coming as Bhārangī were Gardenia latifolia, G. resenifera, G. turgid, Premna intergrifolia and Picrasma quassioides. However, it has been observed that in South Indian Pharmacies, the root and root nodules of Pygmacopremna herbacea were used as Bhārangī in their preparations (K. Vasudevan Nair et.al $(1982)^5$. In a study conducted on the four Indian market samples of Bhārangī, it was found that all samples were adulterated and none of the market sample was found as genuine drug *Clerodendrum serratum*^{3, 4, 9}. Another study showed that the roots of *Bhārangī* were adulterated with the stem of Clerodendrum serratum, mostly in South Indian markets ^{10, 11}. Eventually there were no particular studies conducted on Bhārangī in Kerala markets. This study thus aimed to conclude the exact identity of the market sample of Bhārangī

with special regards to Kerala markets. After undergoing macroscopic examination, morphologically sample 4, 5 & 7 had stout, cylindrical roots similar to genuine root; even then, sample 4 & 7 had a mixture of stem and roots. Sample 2 and 3 possess stoloniferous roots that don't match the original sample ie. Sample 1 which was collected from Mayyanoor, Thrissur.

There were differences in the dimensions and thickness. Sample 5 had similar characters as that of sample 1. Sample 6 was having extremely different morphological characters when compared with the genuine root. Hence, the chances of genuine roots of *Bhārangī* coming in the market is comparatively less based on macroscopical Sample 2 & 3 had similar observations. morphological characteristics of that of roots of Premna herbacea Roxb. In a previous study also stated that sample collected from Southern Kerala can of roots of Premna herbacea Roxb⁴. In-order microscopic character of to compare the Thiruvananthapuram sample, previous study in which the pharmacognostic evaluation of roots of *Pvgmaeopremna* herbacea (syn. Premna herbacea.) discusses the macroscopic, microscopic characteristics along with HPTLC profile of the Premna herbacea Roxbwere noted (Rastogi. S et. al. 2005)¹². None of the samples had similarities with the sample 8 C. indicum 7 .

TS of genuine root of R. serrata was compared with that of 6 market samples. There was extreme difference in case of the Thiruvananthapuram sample (sample 2) and Idukki sample (sample 3) with that of genuine one. Presence of cortical fibers in patches rather than bundle of stone cells. Absence of starch in root samples and the presence of oil globules in both the samples. There was presence of orange-brown colour depositions in the inner cork layers unlike original sample. The Palakkad sample (Sample 4) and Kozhikode sample (sample 5) possess similar microscopic characteristics of that of genuine sample. Palakkad sample possess the stem pieces which did not have the microscopic characteristics of the stem of R. serrata. Presence of oil globules were noted in both the samples. Starch grains were present but less abundant as compared to genuine sample. Wayanad sample (sample 6) and Kasargod sample (Sample7) had no significant similarity with genuine sample. Wayanad sample had well developed xylem resembling a heart wood. Phloem was wide, compressed with brown depositions, fibers are scattered. In Kasargod sample, one of the root pieces showed similar characters of genuine one other sample are different hence it's a mixture of plant parts. None of the samples had similarities with the sample 8 C. indicum.

The HPTLC comparison of water extracts of different samples were carried out. HPTLC chromatograms and densitometric profiles of nine samples at 254nm, 366nm, and 550nm were recorded. The Rf values of the separated compounds of each sample were noted. At 254nm, root of R. serrata had only seven peaks and the stem had eight peaks. The peak intensities were different for each sample, the peak intensities of sample 5 had similarity with that of genuine sample i.e. sample 1. Sample 4 also show some similar peaks but less. Similar Rf values uniformly present in all the samples studied indicate the presence of similar phyto-constituents. Similarly, at 366nm, root of R. serrata had only seven peaks and the stem had nine peaks. Sample 5 shows more similar bands with the sample 1 when compared with others, indicating the presence of a greater number of chemical constituents than the original one. There were no previously published comparable works to conclude these differences in the number of peaks of different samples. After analyzing the result in Table 4, it was clear that the raw drugs available in the market in the name of Cheruthek (Bharangi) were not derived from a single botanical source, revealing the practice of adulteration and substitution in the raw drug industry.

Sl. no.	Samples	Remarks
1	Sample 2- Thiruvananthapuram	Stoloniferous roots- sample had similar characters of roots of Premna herbacea
2	Sample 3- Idukki	Root along with other plant parts it was mostly a mixture. Sample were mixture
		of roots
3	Sample 4- Palakkad	The samples were mixture of various plant parts
4	Sample 5- Kozhikode	The sample had characters of Rotheca serrata
5	Sample 6- Wayanad	The sample was adulterant
6	Sample 7- Kasargod	The sample was a mixture of various plant parts- Adulterant

TABLE 4: INTERPRETATION OF MARKET SAMPLES

CONCLUSION: The samples of the drug *Bhāraṅgī* (*Cheruthek*) collected from 6 different markets from different districts of Kerala were pharmacognostically different, except the sample collected from Kozhikode. The organoleptic and microscopic evaluation of the market samples revealed that only Kozhikode sample had similar characteristics with that of the genuine sample of *Bhāraṅgī- Rotheca serrata* (L.) Steane & Mabb. Even though the genuine part used was root, Kozhikode sample possess both the stem and root. HPTLC profiling also revealed that similar peaks with different band thickness may be due to

differences in maturity, soil, climate, and time of collection of drugs. From the previous reports, the Thiruvananthapuram sample can be the roots of *Premna herbacea* one of the adulterants of the drug *Bharangi*. The Idukki sample too contains roots of *Premna herbacea* along with other mixtures. The Palakkad sample contains the roots and stems of R. serrata and other mixtures. The Wayanad and Kasargod samples were completely different and had no similarity with the source plants *R. serrata & C. indicum*. Hence, it was clear that none of the market samples were only root but a mixture of root and stem.



FIG. 1: SAMPLES OF BHARANGI ALONG WITH SAMPLES FROM DIFFERENT MARKETS IN KERALA. A-B. Genuine sample bharngi; a. root of rotheca. b. stem of *Rotheca serrata*; c. root of *Clerodendrum indicum*; d. Thiruvananthapuram market sample; e. Idukki market sample; f. Palakkad market sample g. Kozhikod market sample, h. Wayanad maekwt sample; i. Kasargod market samole.

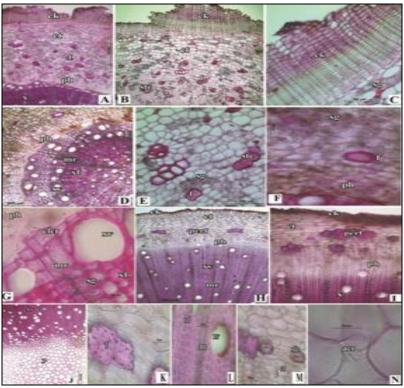


FIG. 2: MICROSCOPIC CHARACTERS OF THE ROOT AND STEM OF R. SERRATA. A-G. TS OF ROOT. A. TS of root portion enlarged; b. TS portion inlarge showing outer portion; c. TS cork region enlarged; d. ts showing stelar region; e. and f. enlarged view of cortical region; g. xylum region portion enlarged; h-n. TS of stem; h. t sod stem portion enlarged; i. TS portion; k. pericycle fibers l. xylume vessels m. enlarged cortical region n. parenchyma cells of pith ck, cork; clcr, crystals of calcium oxalte; ct, cortex; f, fiber; mr. modularly rays; ph phloem; sg starch grains; stc, stone cells; x, xylem; xf, xylem fibers; xv, xylem vessels

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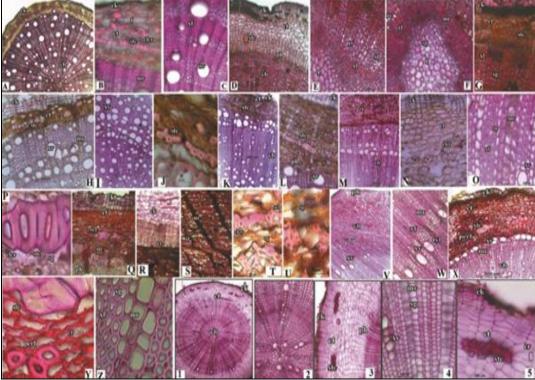


FIG. 3: A-Z: MICROSCOPY OF MARKET SAMPLE OF BHARANGI. A-C. TS of thiruvanthapuram market sample; D-G. TS of Idukki market sample; H-L. TS of palakkad markets saple; M-Q. TS of Kozhikode market sample; R-W. TS of wayanad sample; X-Z. TS of kasargod market sample; 1-5 microscopy *of Clerodendrum indicum*.

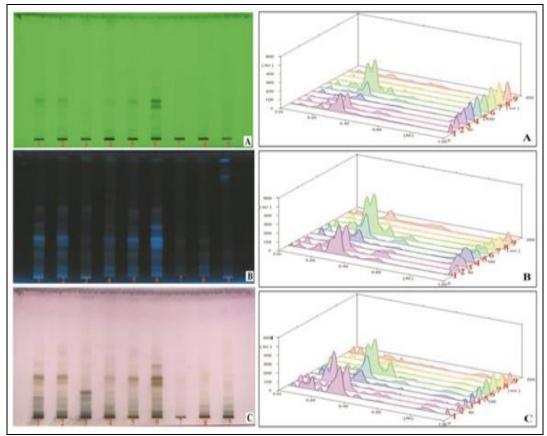


FIG. 4: HPTLC COMPARISON OF GENUINE AND MARKET SAMPLES OF BHARANGI. A-c. TLC profile of sample at 254nm, 366nm after derivatisation; d-f; denitometeric profile at 254nm 366nm and after derivatisation. 1. *Rotheca serrata* rot, 2.Rotheca serrata stem, 3. Thiruvananthapuram sample, 4. Idikki sample, 5. Palakkad sample, 6. Kozhikode sample, 7. Wayanad sample, 8. Kasargod sample, 9. Root of *Clerodendrum indium*.

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ACKNOWLEDGEMENT: The authors are grateful to Dr. N. Manoj Kumar, Prof. and Head; Dr. Madhu K P Associate Prof., Dr. Vidya Unnikrishnan, & Dr. Jyolsna G Krishna, Assistant Prof.; Department of Dravyaguna Vijnana, VPSV *Ayurveda* College, Kottakkal; Mr. Deepak M, Scientist, Phytochemistry Division & Ms. Haritha, Technical Assistant, Pharmacognosy Division, CMPR, AVS, Kottakkal, Kerala for their help and support during the study.

Financial Support and Sponsorship: Nil

CONFLICTS OF INTEREST: There was no conflict of interest.

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How to cite this article:

Navami A, Vivek P and Harinarayanan CM: Pharmacognostic and HPTLC comparison of Kerala market samples of Bharangi (Cheruthek) with genuine source plant *Rotheca serrata* (L.) steane & MABB. Int J Pharm Sci & Res 2023; 14(12): 5676-86. doi: 10.13040/IJPSR.0975-8232.14(12).5676-86.

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