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PHARMACOGNOSTICAL STANDARDIZATION ON THE ROOTS OF AGNIMANTHA - *PREMNA SERRATIFOLIA* LINN.

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ABSTRACT: Present communication deals with the pharmacognostical standardization on the roots of *Agnimantha* (*Premna serratifolia* L. (syn. *P. integrifolia* L.) belong to the family Verbenaceae. It is an important medicinal plant used in many Ayurvedic formulations. According to Ayurveda the roots and leaves are most useful parts of the plant. It is one among the *Dashamula*. Macro, microscopical and physicochemical studies were carried out as per the latest Ayurvedic Pharmacopoeia of India, part-1, volume-8 protocols to establish its Pharmacognostical standards. Studies revealed the presence of abundant, simple to compound starch grains with prominent hilum in the centre, cork cells filled with yellowish brown pigmented cells, starch grains in the medullary rays, abundant xylem vessels with bordered pits and tracheids with pits, prism shaped crystals in the cortex region. Phytochemical studies showed the presence of carbohydrates, alkaloids, phenols, tannins etc.

INTRODUCTION: Agnimantha a sanskrit name equivalent to *Premna serratifolia* Linn. (Synonym *P. integrifolia* Linn.) is a large, thorny deciduous shrub or a tree, up to 9 meters in height, common along the Indian and the Andamans coasts, it also occurs in the plains of Assam and in Khasi hills. In Ayurveda the roots and leaves are most used parts for many ailments. The roots are considered as astringent, stimulant, liver tonic, laxative, carminative and antibacterial. Leaves infusion with pepper is useful in cold and fever, leaf sap is useful to alleviate headache^{1,2}.

Roots are useful in vitiated conditions of *vata*, *kapha*, neuralgia, inflammations, cardiac disorders, cough, asthma, bronchitis, leprosy, skin disorders, dyspepsia, flatulence, constipation, fever, diabetes and anorexia etc.³ In Vietnamese traditional medicine, the leaves of *S. chinensis* are used to treat indigestion, dysuria and dysentery; roots against indigestion, stomach ache and fever⁴. The root methanolic extract showed better immunomodulatory activity in experimental mice⁵. A paste of its roots with water and butter fat is given in Urticaria⁶.

Ayurvedic Description of Properties: In terms of Rsasapanchaka theory of Ayurveda, the properties are as follows: Rasa (taste) - Tikta (bitter), kaṭu (pungent), kaṣāya (astringent); Guṇa (quality) - Rūkṣa (creates dryness), laghu (light for digestion); Vīrya (potency) - Uṣṇa (Hot in potency); Vipāka (digestive effect) - Kaṭu (pungent); Karma (action)

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- śothahara (Anti inflammatory), vedanāsthāpana (anodyne or which relieves pain); Doṣāghnatā (effect on doshas) - Kapha-vātahara (Acts against the diseases came due to *Kapha dosha* and *Vata dosha*)⁷.

As per the 3 volumes of Ayurvedic Formulary of India, more than 40 formulations has been prepared by using Agnimantha as one of the ingredient, some of them are as follows: *Amrtarista, Dantyadyarista, Dasamularista, Agastya Haritaki Rasayana, Cyavanaprasa, Brahma Rasayana, Bharnigiguda, Dasamula Katutrayakavthacurna, Dasamulakvathacurna, Dasamulapancakoladikavthacurna, Darunagaradikavthacurna, Rasnadikavthacurna, Saptasarakavthacurna, Indukanta Ghrta, Dasamulaghrta, Dasamulasaptalakaghrta, Dadhikaghrta, Dhanvantaraghrta, Mahapancagavyaghrta, Lasunadighrta, Sukumaraghrta, Dhanvantarataila, Narayanataila, Prabhanjana Vimardhanataila, Brha-tmasataila, Mahanarayanataila, Mahavisagarbhataila, Sahacaraditaila*⁸; *Brhacchagaladyaghrta, Guducy-aditaila, Dasamulataila, Madhyama Narayanataila, Musikadyataila, Vajrakapata rasa, Vatagankusa rasa, Sirahsuladivajra rasa*⁹; *Nalikerasava, Dasamularka, Vasisthaharita-kyavaleha, Caturda-sanga-kvathacurna, Tagaradikvathacurna, Bhunimbadi Astadasangakavthacurna, Viratarvadiganakavtha-curna etc*¹⁰.

Chemical Constituents: Premnacorymboside A, scutellarioside II, quercetin - 3-rutinoside, leonurioside A were isolated from the stem bark⁴. Alkaloid contents like premnine, ganikarine, premnazole are reported in root from different researchers⁵. 1β, 3α, 8β-trihydroxy-pimara-15-ene; 6α, 11, 12, 16-tetrahydroxy-7-oxo-abieta-8, 1, 13-triene; 2α, 19-dihydroxypri-mara-7,15-diene were identified in root bark¹¹. It is known as follows in different Regional languages in India.

Sanskrit: Agnimantha, Ganiparnika, Jjaya, Vaijayanti, Arani

Bengali: Bhut-bhiravi

Gujarathi: Mothi-arni

Hindi: Agetha

Kannada: Agnimanda, Eegigida

Malayalam: Munna

Tamil: Munnay

Telugu: Pomantai, Peddanarva, Gaebunelli;

Urdu: Arani¹.

MATERIALS AND METHODS: The roots of Agnimantha were collected from Budnal village, Huballi taluk, Dharwad district, Karnataka, authenticated by Survey of medicinal plant unit, Regional Ayurveda Research Institute for Metabolic Disorders, Bangalore. The roots were shade dried, some are pulverized by mechanical grinder to get coarse powder and stored in a closed vessel, to carryout microscopical studies, powder studies, physicochemical and preliminary phytochemical analysis as per the standard protocols. Macroscopical, microscopical and powder studies were carried out as per the standard procedures^{12, 13}.

Hydro-Alcoholic Extract: 100 g of coarse powder was taken in an extractor and mixed with 50% aqueous alcohol, thrice the quantity of raw material and refluxed for 3 - 4 h. The extract was filtered through a whatmann no.1 filter paper. The process was repeated three times and each time the extract was filtered into the same vessel. The combined filtrates were concentrated to syrupy consistency and dried by using a rotary evaporator¹⁴.

Water Extract: 100g of coarse powder was taken in an extractor and was added with thrice quantity of distilled water and heated at a temperature 80 °C for 3 - 4 h. The extract was filtered through a whatmann no.1 filter paper. The process was repeated three times and each time the extract was filtered into the same vessel. The combined filtrates were concentrated to syrupy consistency and dried using a rotary evaporator¹⁴.

Thin Layer Chromatography: Shade dried roots coarse powder was extracted with methanol, water-alcohol and water by reflux method. TLC studies of these extracts were carried out by using, commercially available precoated plates at room temperature by following standard procedures¹⁴.

Physicochemical Analysis: The powdered drug was used to carry out various physicochemical parameters such as total ash, acid-insoluble ash, loss on drying at 105 °C, water soluble and alcohol soluble extractive values. The water and water-alcoholic extracts were dried and subjected to various tests like total ash, acid-insoluble ash, pH of aqueous solution and total soluble solids as per the latest Ayurvedic Pharmacopoeia of India protocols¹⁴.

Preliminary Phytochemical Analysis: The powdered drug was extracted with water, alcohol and petroleum ether and methanol for carrying out different preliminary phytochemical tests for alkaloids, carbohydrates, proteins, phenols, tannins, saponins, starch, flavonoids and steroids etc. by using Standard testing protocols^{15, 16}.

RESULTS:

Macroscopic Characteristics: Roots light brown or yellowish brown, woody, 6 - 8 cm long and 4 - 5

mm in diameter, brown from outside and yellow from within, having a short fracture. Roots contain yellow pigment tannin. Roots are light brown, woody, branched and cylindrical. Outer surface is exfoliated at some places and shows longitudinal striations and wrinkles. Roots possess bland taste and are slightly aromatic, woody, branched and tortuous to cylindrical in shape. Surface gets exfoliated easily and shows prominent longitudinal striations and wrinkles **Fig. 1** and **2**.



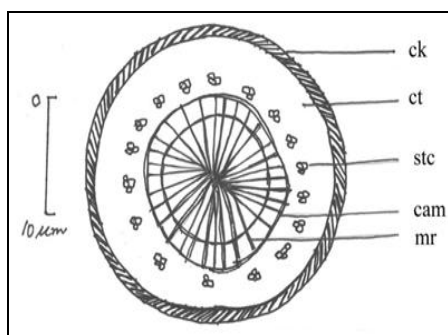
FIG. 1: LIVE PLANT



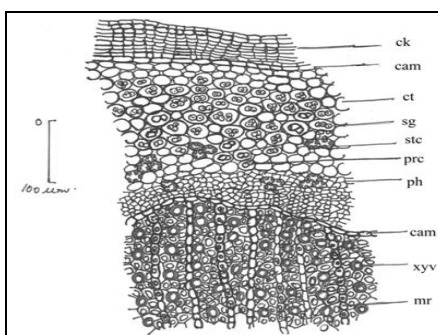
FIG. 2: DRIED ROOT PIECES

Microscopic Characteristics: The transverse section shows cork, consisting of 8 - 10 rows of tangentially elongated and radially arranged suberised cells. Cork is followed by cortex made up of 10 - 15 layers of thin walled parenchymatous cells, closely arranged with little intercellular spaces. Cortex shows two to three discontinuous layers of stone cell that are thick walled, lignified and lodged with prisms of calcium oxalate.

A few cortical cells contain yellowish brown pigment. Phloem shows thin walled cells and in between the phloem cells stone cells with prisms of calcium oxalate crystals are present. The xylem vessels are of varying size, lignified, found isolated or in the group of 2 - 3. Medullary rays are 2 - 3 seriate and the cells are pitted and lignified. Starch is present in the medullary ray cells **Fig. 3**.



T. S. OF ROOT (DAIGRAMMATIC)



T. S. OF ROOT

FIG. 3: MICROSCOPY OF PREMNA SERRATIFOLIA LINN.

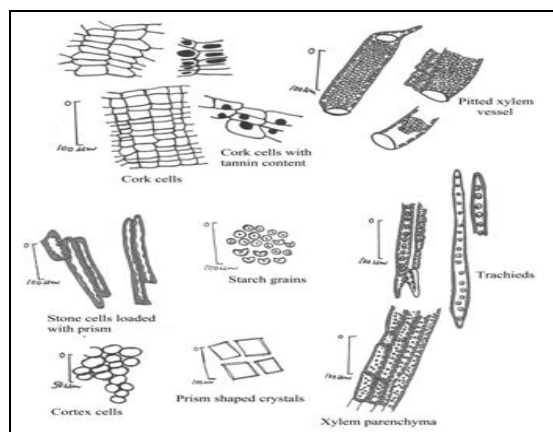
Ck: cork, Cam: cambium, ct: cortex, mr: medullary ray, ph: phloem, prc: prismatic crystal of calcium oxalate, stc: stone cell, sg: starch grains, xyv: xylem vessel

Powder Microscopy: Powdered root is light brown, with slightly aromatic and bland taste. Starch is abundant, simple, spherical and cup shaped. Stone cells are rectangular to oblong and lodged/ loaded with 3 - 8 prisms of calcium oxalate

which are found scattered also. The vessels and fibers of the xylem are lignified and found in the groups of interlocking cells. The xylem vessels shows bordered pitted thickenings **Fig. 4**.

Diagnostic Characters:

- Presence of abundant starch grains in the cortex region.
- Presence of prism shaped crystals and stone cells in the cortex region.
- Presence of yellowish brown pigmented cells in the cork region.
- Presence of starch grains in the medullary ray cells.
- Presence of abundant xylem vessels with bordered pits.

FIG. 4: POWDER STUDY OF *PREMNA SERRATIFOLIA* L.

Physicochemical Analysis: Physicochemical parameters such as total ash, acid-insoluble ash, loss on drying at 105 °C, water soluble and water-alcohol soluble extractive values were carried out and results were given in **Table 1**. The water and water-alcoholic extracts were dried and subjected to various tests like total ash, acid-insoluble ash, pH of aqueous solution and total soluble solids as per the latest Ayurvedic Pharmacopoeia of India protocols and results were given in **Table 2**.

Preliminary Phytochemical Analysis: Preliminary phytochemical analysis have been carried out for alkaloids, carbohydrates, proteins, phenols, tannins, saponins, starch, flavonoids and steroids *etc* and the results were given in **Table 3**.

TABLE 1: PHYSICOCHEMICAL PARAMETERS *P. INTEGRIFOLIA* LINN. ROOT POWDER

Name of the parameter	Values (%) w/w
Description	Light brown coarse powder
Loss on drying at 105 °C	6.23
Total ash	10.18
Acid-insoluble ash	1.25
Water-soluble extractive	15.79
Alcohol-soluble extractive	6.05
Water-Alcohol (50:50) soluble extractive	9.86

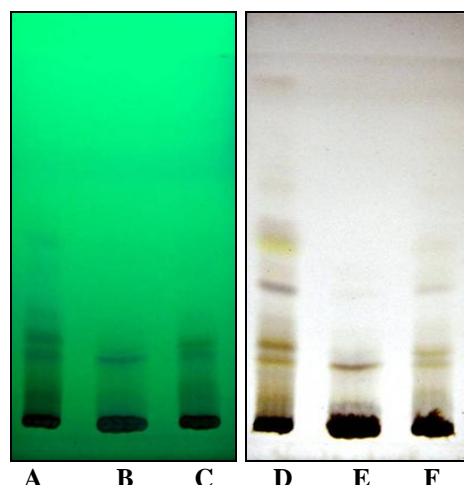
TABLE 2: PHYSICOCHEMICAL PARAMETERS OF *P. SERRATIFOLIA* L. ROOT POWDER EXTRACTS

Name of the parameter	Values (%) w/w	
	Hydro-alcoholic extract	Water extract
pH (5 % w/v aq. solution)	5.05	4.85
Loss on drying at 105 °C	9.38	10.86
Total ash	5.97	13.28
Acid-insoluble ash	0.25	0.20
Total soluble solids	95.39	92.93

TABLE 3: PRELIMINARY PHYTOCHEMICAL TESTS FOR DIFFERENT EXTRACTS OF *P. SERRATIFOLIA* L. ROOT

Natural product group	Test for natural products	Presence (+)/Absence (-)
Alkaloids	Dragendorff's test	+
	Hager's test	+
	Mayers's test	+
	Wagner's test	+
	Anthrone test	++
Carbohydrates	Benedict's test	++
	Fehling's test	++
	Molisch's test	++
	Flavonoids	--
Phenols	Ferric chloride test	+
	Lead acetate test	+
Proteins	Biuret's test	+
	Millon's test	+
Saponins	--	+
Steroids	Salkowski reaction	-
	Tannins	Ferric chloride test
	Lead acetate test	+

Thin Layer Chromatography: Thin layer chromatography was carried out on a precoated silica gel 60₂₅₄ plate. Few drops of test solution applied on a TLC plate as band. Developed the plate to a distance of 8 cm from the line of application by using Hexane: Ethyl formate (8:2) as mobile phase (solvent system).

**FIG. 5: THIN LAYER CHROMATOGRAPHY**

A: Powder ext., B: water ext., C: Wat.-Alc. ext. under UV254 nm; D: Powder ext., E: water ext., F: Wat.-Alc. ext. after spraying with 10% H₂SO₄).

Dried the plate in air and examined under 254 nm, recorded the R_f values. Sprayed the TLC plate with 10% sulphuric acid reagent. Heated the plate at 110 °C for about 5 min and recorded the R_f values. The chromatogram obtained with powder test solution shows major R_f values under UV-254nm ~ 0.06, 0.175, 0.21, 0.30, 0.56 and after spraying at ~ 0.06, 0.175, 0.21, 0.30, 0.375, 0.50, 0.56, 0.93. The hydro-alcoholic extract test solution shows major R_f values under UV-254 nm at ~ 0.06, 0.175, 0.21 and after spraying at ~ 0.06, 0.175, 0.21, 0.375. Water extract test solution shows major R_f values under UV-254 nm at ~ 0.21 and after spraying at ~ 0.06, 0.21 **Fig. 5**.

CONCLUSION: Agnimantha is an important Ayurvedic drug used in several Ayurvedic preparations. Studies carried out on the roots of Agnimantha - *Premna integrifolia* Linn. helps in the identification/authentication of different species of *P. integrifolia* and also different sources of Agnimantha. Since agnimantha is also used under different botanical names like *Clerodendrum phlomidis* Linn. f., belonging to the same family Verbenaceae, this type of studies helps in differentiating the other sources of agnimantha, which will be more useful for researchers, students and Ayurvedic Physicians for identifying the correct drug and it may also be included in the Ayurvedic Pharmacopoeia of India.

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CONFLICT OF INTEREST: Nil

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