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PHARMACOGNOSTICAL AND PHYTOCHEMICAL PROFILES OF Peltophorum pterocarpum LEAVES

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ABSTRACT

Peltophorum pterocarpum (Family- Caesalpiniaceae) is used in different systems of traditional medicine. The present study was planned to investigate the macroscopy, microscopical characters, quantitative profiles, powder microscopy, physicochemical parameters of leaves of *Peltophorum pterocarpum* and preliminary phytochemical screening of HAEPP (Hydroalcoholic extract of *Peltophorum pterocarpum*). Pharmacognostical parameters are helpful in identification, authentication and control the adulteration of crude drugs. The preliminary phytochemical screening reveals the presence of alkaloids, carbohydrates, flavonoids, proteins, amino acids, sterols, terpenoids, tannins and coumarins.

Key words: Peltophorum pterocarpum, Traditional medicine, Pharmacognosy, phytochemistry.

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INTRODUCTION

In many communities around the world, plants are the primary source of health care. They also serve as resources for the development of biomedicine and traditionally used species have made significant contributions to the advancement of biomedical drug development (Estevão N. F., 20118). *Peltophorum pterocarpum* (Family- Caesalpiniaceae) is a large ornamental tree grown around the world. This plant is commonly known as copper-pod, yellow Poinciana and golden flamboyant. Tamil name is perungondrai and iyalvagai.

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deciduous tree has a rounded canopy and is capable of reaching 50 feet in height with a 30 to 50-foot spread. *Peltophorum pterocarpum* is the important plant species in traditional medicine (Dharmasoth Rama Devi and Ganga Rao Battu., 2018, Arul Sheeba Rani, M and Mary Josephine, R., 2018). The plant is used in different parts of the world for the treatment of several ailments like stomatitis, insomnia, skin troubles, constipation, ringworm, dysentery, muscular pains, sores, and skin disorders (Corners, E. J. H., 1997, Tan. Hugh T.W. and T. Morgany., 2001, Nathan VK., 2012, Swee Ping and Wee Mei Lynn., 2001).

This upright, handsome, spreading, semi-evergreen,

MATERIALS AND METHODS

PLANT COLLECTION AND AUTHENTICATION

Leaves of *Peltophorum pterocarpum* was collected from Alwarpuram, Madurai district and it was authenticated by **Dr. D.Stephen, M.Sc., Ph.D.,** Assistant Professor, Department of Botany, The American College, Madurai -20, Tamil Nadu. The herbarium of this specimen was kept in the department for further reference.

PHARMACOGNOSTICAL STUDIES MACROSCOPICAL STUDIES

The fresh leaves are studied for its morphological characters like colour, odour, taste, shape, arrangement, apex, base, margin, length and width by organoleptic evaluation.

MICROSCOPICAL STUDIES TRANSVERSE SECTION

Sample (fresh *Peltophorum pterocarpum* leaves) was preserved in fixative FAA (Formalin -5ml + Acetic acid -5ml +70% Ethyl alcohol -90ml) for more than 48 hr. The preserved specimens were cut into thin transverse section using a sharp blade and the sections were stained with safranin. Transverse sections were photographed using trinocular microscope under bright field light.

QUANTITATIVE MICROSCOPY

The vein islet and vein termination number, stomatal number and stomatal index were determined on fresh leaves by using standard procedure.

POWDER MICROSCOPY

A pinch of the powdered sample was mounted on a microscopic slide with a drop of 50% glycerol. Characters were observed using trinocular microscope and Photomicrographs of diagnostic characters were captured and documented (Iyengar M.A and Nayak S.G.K., 1994, Wallis TE., 1946).

PHYSICO-CHEMICAL PARAMETERS

The foreign matters, loss on drying, extractive value and ash value were determined according to the methods given in pharmacopoeia (Anonymous., 1998, Anonymous., 2001).

PHYTOCHEMICAL STUDIES EXTRACTION

The leaves of *Peltophorum pterocarpum* was collected, shade dried and coarsely powdered. The powdered leaves were defatted with petroleum ether and then the defatted marc was allowed to macerate with hydroalcohol using 70% ethanol for about 72 hours. The extracts were filtered through whatman filter paper No. 42 (125 mm). The entire extracts were concentrated to dryness. The final dried extract was stored in an air tight and labelled container.

PRELIMINARY PHYTOCHEMICAL ANALYSIS

Various phytochemical tests were performed to identify the phytoconstituents present in HAEPP by using standard protocol (Harbone JB., 1973, Kokate C.K., 1994., Payal M. Panchal., 2012, Abdullahi R., 2020).

RESULTS AND DISCUSSION MACROSCOPICAL STUDIES

Macroscopy of the *Peltophorum pterocarpum* leaves was studied (Figure 1) and reported (Table-1). The leaves of *Peltophorum pterocarpum* is bipinnate, alternate, stipulate, dried leaves are yellowish green in colour. Odour and taste are not characteristic.

TS OF RACHILLA

TS of rachilla shows Single layered epidermis covered with thick cuticle shows uni to multi cellular covering trichomes. Ground tissue possesses centrally arranged prominent vascular bundle and two trace bundles located at the upper side, under each elevation. Vascular bundles are closed; phloem consists of usual elements embedded with a few rosette crystals of calcium oxalate. Xylem consists of radially arranged vessels surrounded by tracheids and fibres. In the centre small parenchymatous pith is found (Figure 2- 5).

Leaflets

Midrib

TS of leaflet passing through midrib are broadly convex at the lower side and shows slightly convex upper surface. Single layered epidermis covered with thick cuticle is followed by a sub-epidermal layer having one to two layers of parenchymatous cells over the vascular bundle. Centrally placed vascular bundle is closed and collateral (Figure 6).

Lamina

TS of lamina are dorsiventral, hypostomatic, shows single layered upper and lower epidermis covered with thick cuticle and simple covering trichomes (Figure 7, 8).

QUANTITATIVE MICROSCOPY

The quantitative parameters of leaves of *Peltophorum pterocarpum* were recorded (Table 2). The leaf lower side shows numerous paracytic, few anomocytic stomata and they are very rare on the upper surface (Figure 9-11).

POWDER MICROSCOPY

Peltophorum pterocarpum leaves powder was brownish green in colour and shows various powder characters (Figure 12).

PHYSICO-CHEMICAL PARAMETERS

Foreign matter, loss on drying, extractive value and ash values were determined and reported (Table 3).

PRELIMINARY PHYTOCHEMICAL ANALYSIS

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The preliminary phytochemical screening reveals the presence of alkaloids, carbohydrates, flavonoids, proteins, amino acids, sterols, terpenoids, tannins and coumarins (Table 4).

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TABLE: 1 MORPHOLOGICAL FEATURES OF Peltophorum pterocarpum LEAF LEAE TYPE Bisingetely compound

LEAF IYPE	Bipinnately compound
LEAF ARRANGEMENT	Alternate, 11- 17 pairs of leaflets.
LEAF SHAPE	Oblong
SIZE	Rachis 25 to 35 cm long
	Lamina 1 to 2 cm length and 0.4 to 0.8 cm width
LEAF VENATION	Pinnate
LEAF – MARGIN	Entire
APEX	Obtuse to retuse
BASE	Truncate
BENEATH	Puberulent
MIDRIB	Prominent
LEAF COLOUR	Fresh leaf – Green
	Dried leaf – Yellowish green

TABLE: 2 QUANTITATIVE MICROSCOPY OF Peltophorum pterocarpum LEAF

PARAMETERS	UPPER EPIDERMIS (/mm ²)	LOWER EPIDERMIS (/mm ²)
Epidermal number	1536 - 1648	1344 - 1616
Stomatal number	-	208 - 256
Stomatal index	-	13.90
Palisade ratio	4-7	
Vein islets number	9	
Vein termination number	31	

TABLE: 3 PHYSICO-CHEMICAL PARAMETERS OF Peltophorum pterocarpum LEAF

S. NO	PARAMETERS	RESULTS
1	FOREIGN MATTER	NIL
2	LOSS ON DRYING	$10.88 \pm 0.126 \ \% w/w$
3	EXTRACTIVE VALUE	
	 Petroleum ether extract 	1.46 ± 0.042 %w/w
	 Chloroform extract 	$4.06 \pm 0.117 \ \% w/w$
	 Acetone extract 	7.53 ±0 .218 %w/w
	 Ethanolic extract 	$13.20 \pm 0.381 \ \text{\% w/w}$
	 Hydroalcoholic extract 	$23.06 \pm 0.665 \ \% \text{w/w}$
	 Diethyl ether extract 	$1.86 \pm 0.054 \ \% w/w$
	 Aqueous extract 	$12.40 \pm 0.357 \ \text{\% w/w}$
4	ASH VALUE	
	✤ Total ash	$11.18 \pm 0.129 \ \% w/w$
	✤ Water soluble ash	$5.66 \pm 0.065\% w/w$
	 Acid insoluble ash 	$9.26 \pm 0.107 \ \text{\% w/w}$

TABLE: 4 PRELIMINARY PHYTOCHEMICAL ANALYSIS OF HAEPP

S.NO	TEST	INFERENCE
1	ALKALOIDS	PRESENT
	MAYER'S TEST	+
	WAGNER'S TEST	+
	HAGER'S TEST	+
	DRAGENDORFF'S TEST	+

2	CARBOHYDRATES	PRESENT
	MOLISCH'S TEST	+
	FEHLING'S TEST	+
	BENIDICT'S TEST	+
3	GLYCOSIDES	
	ANTHROQUINONE GLYCOSIDES	ABSENT
	BORNTRAGER'S TEST	-
	MODIFIED BORNTRAGER'S TEST	-
	CARDIAC GLYCOSIDES	ABSENT
	KELLER KILIANI TEST	-
	LEGAL'S TEST	-
	BALGET'S TEST	-
4	STEROLS	PRESENT
	SALKOWSKI'S TEST	+
	LIBERMAN BURCHARD'S TEST	+
5	FLAVONOIDS	PRESENT
	SHINODA TEST	+
	ALKALI TEST	+
	LEAD ACETATE TEST	+
	ZINC HYDROCHLORIDE TEST	+
6	PROTEINS	PRESENT
	MILLON'S TEST	+
	BIURET TEST	+
7	AMINO ACIDS	PRESENT
	NINHYDRIN TEST	+
	NITRIC ACID TEST	+
8	TERPENOIDS	PRESENT
	NOLLER'S TEST	+
	TRITERPENOIDS	PRESENT
9	GUM	ABSENT
10	MUCILAGE	ABSENT
11	SAPONINS	ABSENT
	FOAM TEST	-
12	TANNINS	PRESENT
	FERRIC CHLORIDE TEST	+
	GELATIN TEST	+
13	VOLATILE OIL	ABSENT
14	FIXED OIL AND FATS	ABSENT
15	COUMARINS	PRESENT

FIGURE: 1 Peltophorum pterocarpum leaf







FIGURE: 3 TS OF RACHILLA UPPER PORTION ENLARGED VIEW



FIGURE: 4 TS OF RACHILLA MIDDLE PORTION ENLARGED VIEW



FIGURE: 5 TS OF RACHILLA LOWER PORTION ENLARGED VIEW



Col - collenchyma; Cu - cuticle; E - epidermis; MR - medullary ray; Pa - parenchyma; Per - pericycle; Ph - phloem; RCr - rosette crystal; T - trichome; TB - trace bundle; Ve - vessel; VB - vascular bundle

FIGURE: 6 TS OF Peltophorum pterocarpum MIDRIB



FIGURE: 7 TS OF LAMINA PASSING THROUGH MIDRIB



FIGURE: 8 TS OF LAMINA PASSING THROUGH MARGIN



 $\begin{array}{l} Cu-cuticle; \ \overline{E-epidermis}; \ Lat-laticiferous \ cells; \ LE-lower \ epidermis; \ Me-mesophyll; \ Pa-parenchyma; \ Pal-palisade; \ Per-pericycle; \ Ph-phloem; \ SE-sub-epidermis; \ Sp-spongy \ parenchyma; \ T-trichome; \ UE-upper \ epidermis; \ Ve-vein; \ Xy-xylem \end{array}$

FIGURE: 9 VEIN ISLET AND VEIN TERMINATION



FIGURE: 10 UPPER EPIDERMIS



FIGURE: 11 LOWER EPIDERMIS



Cic – cicatrix; E - epidermis; St - stomata; VT - vein islet; VI - vein termination







Pitted parenchyma



CONCLUSION

The Pharmacognostical parameters of *Peltophorum pterocarpum* leaves are helpful in identification, authentication and acquire genuine plant materials. The phytochemical studies revealed the presence of various phytoconstituents. These parameters are very helpful in the new drug development.

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