

Research Journal of Pharmaceutical, Biological and Chemical Sciences

REVIEW ARTICLE

Ethno-medicinal, Phytochemical and Pharmacological review of an amazing medicinal herb *Peperomia pellucida* (L.) HBK.

Pulak Majumder^{1*}, Priya Abraham¹, Satya V²

¹Dept. of Pharmacognosy, Rajiv Gandhi Institute of Pharmacy, Trikaripur, Kasaragod (Dist.), Kerala-671310. ²Dept. of Pharmaceutical chemistry, Rajiv Gandhi Institute of Pharmacy, Trikaripur, Kasaragod (Dist.), Kerala-671310.

ABSTRACT

Peperomia pellucida (L.) HBK (Fam. Piperaceae) has been used as a (Rasayan) drug in the Ayurvedic system of medicines. Peperomia pellucida is reported to posse antipyretic, analgesic, anti-inflammatory, antimicrobial, refrigerant and CNS activity. Traditionally it is used in the treatment of headache, fever, eczema, abdominal pains, and convulsions. In traditional Ayurvedic system of medicine in India Peperomia pellucida is used as Rasa, Guna and Virya. Phytochemical screening of this plant has shown the presence of flavonoids, tannins, alkaloids, steriod and triterpenoid. Isolation of antifungal and anticancer constituents from this plant was also reported newly. The present study is based on the work done till date regarding the phytoconstituents and pharmacological activity of Peperomia pellucida.

Keywords: Piperaceae, Essential oils, Analgesics, Anti-pyretics, CNS activity.

*Corresponding author E-mail: pulak2007@gmail.com

October - December 2011 RJPBCS Volume 2 Issue 4 Page No. 358



INTRODUCTION

Peperomia pellucida (L.) HBK is also known as shiny bush or silver bush belonging to family Piperaceae. In Sanskrit, it is known as Toyakandha, Varshabhoo. Peperomia pellucida is an herbaceous plant found in many South American and Asian countries. The species develops during rainy periods (often in the spring) and thrives in loose, humid soils under the shade of trees. [1-4]. It grows in moist habitat and is found throughout the major parts of India. In different parts of India it is known with different names like Lochi pata in Bangali [5], Mashitandu chedi in Malayalam and Pononoa in Assamese etc [5,6]. Whole plant or parts of plant are used for different purposes. Despite its wide range of folk medicinal uses in India subcontinent, there is very little scientific documentation available on its pharmacological and biological activities as well as its chemical constituents.

MATERIAL AND METHODS

TAXONOMICAL CLASSIFICATION [7].

Kingdom **Plants**

Subkingdom *Tracheobionta* – Vascular plants Superdivision Spermatophyta – Seed plants Division Magnoliophyta – Flowering plants Magnoliopsida – Dicotyledons Class

Subclass Magnoliidae Order **Piperales** Family Piperaceae Genus peperomia

Species Peperomia pellucida

Botanical description

Peperomia pellucida (L.) HBK is an annular herb. The roots are fibrous; stems translucent pale green, erect or ascending, usually 15-45 cm long, internodes usually 3-8 cm long, glabrous and the leaves are medium green on upper surface, lower surface whitish green, thinly fleshy, drying papery, broadly ovate, 1.5-4 (-5) cm long, 1-3.3 cm wide, palmately 3-nerved or 5nerved, glabrous, apex acuminate, base subcordate to truncate, petioles 0.5-2 (-3) cm long, glabrous. One to several spikes are available, terminal and axillary or leaf-opposed, filiform, ca. 3-6 cm long, the rachis ca. 0.4-0.6 mm in diameter, glabrous, flowers well-spaced, peduncles ca. 0.6-1 cm long, glabrous; ovary ovoid; stigmas terminal and also fruits were subglobose, ca. 0.5 mm long, longitudinally ridged, apex beaked [8].

October - December Volume 2 Issue 4 **RIPBCS Page No. 359**

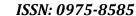


GEOGRAPHICAL DISTRIBUTION

Peperomia pellucida is widely distributed in many South American and Asian countries [4, 9-11]. The plant is occasionally cultivated and sparingly naturalized in Hawaii [8]. In Fiji, this plant is occurring at elevations of sea level to about 400 m as a weed along roadsides, in plantations, on damp ground in shady places near houses, and occasionally along forest trails [12]. Some other region across the glove like southern regions of China [13], Samoa, New Guinea etc are the places where this plant are found in damp shady conditions and living bare ground which is mostly from sea level to 700 m [14,15]. The plant mostly found at tropical and subtropical parts of India. Basically the Native range of this plant is Tropical America, but now widely cultivated and naturalized throughout the tropics [8].

TRADITIONAL USES

Plants have played a significant role in maintaining human health and improving the quality of human life for thousands of years, and served as valuable components of food and medicines [16]. Peperomia pellucida leaves and stems may also be eaten as vegetable [17]. In salads, the fresh plant has the crispness of carrot sticks and celery. As Ethno-medicinal uses of this plant *Peperomia pellucida* has been applied for treating abdominal pain, abscesses, acne, boils, colic, fatigue, gout, headache, renal disorders, and rheumatic joint pain[6,18,19,20]. In Bolivia, Altenos Indians use the whole plant to stop hemorrhages. The roots are used to treat fevers and the aerial parts are used as dressing for wounds [21]. In northeastern Brazil, the plant has been used to lower the cholesterol level [9]. In Guyana and the Amazon region, it is a popular cough suppressant, emollient, and diuretic. It is also used to treat proteinuria [3, 4]. In the Philippines, a decoction of the plant is used to decrease uric acid levels and to treat renal problems [5]. In different region of Lakshmipur district of Bangladesh, the leaves of the plant are used by local people in the treatment of excited mental disorder [22]. It is also used topically for skin disorders such as acne and boils. In South America, A solution of the fresh juice of stem and leaves is used against eye inflammation [6]. Infusion and decoction of leaves and stems are used for gout and arthritis. According to Manila Medical Society P. pellucida is used to relieve arthritic pains, but can cause CNS depression [23]. This plant has externally used as a facial rinse for complexion problems. Pounded whole plant used as warm poultice for boils, pustules and pimples and also used for headaches, rheumatic pains and impotence [24]. Peperomia pellucida is also used in traditional Ayurvedic medicine [6]. It is described in Ayurveda as – Rasa – Katu and Madhur; Guna- Lakhu, rooksha, Teekshna; and Virya- Ushna. The plant is described to passify vitiated cough, pitta, constipation, kidney diseases, urinary retention, dysuria, urinary tract infections, emaciation, edema and general weakness. Infusion and decoction of leaves and stems of fresh plant are eaten as salad for the treatment of gout and arthritis [24, 25]. According to Ethno-botanical studies the whole plant has been in medicinal use since long. It is crushed and mixed with water to form a mixture, heated and administered orally to cure hemorrhage. It is also been applied against coughing, fever, common cold, headache, sore throat, diarrhea, against kidney and prostate problems and against high blood pressure [26].





PHYTOCONSTUENTS

The plant *Peperomia pellucida* was found to have variety of chemical constituents. Phytochemical screening revealed the presence of alkaloids, cardenolides, saponins and tannins, while anthraquinones was observed to be absent [27]. Stem also contain alkaloid, tannins, flavanoids and steriods, except saponins. The roots of *Peperomia pellucida* also had shown the presence of alkaloid, tannins, steroids and carbohydrates etc. The essential oils of the plant were found primarily in medical literature. One study identified 71 compounds from the essential oils of 10 Piperaceae species. Sesquiterpenes appear to be the major chemical constituents in the essential oils. Carotol (13.41%) was the major hydroxylated sesquiterpene in a chemical analysis of *Peperomia pellucida*. Flavonoids, phytosterols, arylpropanoids (eg, apiols), substituted styrenes, and a dimeric ArC₂ compound or pellucidin A have been isolated. Antifungal activity has been documented for arylpropanoids such as the apiols. Other compounds, like the peperomins, have cytotoxic or anticancer activity in vitro. Isolated flavonoids include acacetin, apigenin, isovitexin, and pellucidatin. Isolated phytosterols include campesterol and stigmasterol.

Also contains five new compounds (1-5), including two secolignans, two tetrahydrofuran lignans, and one highly methoxylated dihydronaphthalenone. These compounds were accompanied by the known peperomins A, B, C [28], and E [29], 7, 8 - trans 8, 8' – trans -7', 8'- cis- 7, 7' – bis (5-methoxy-3,4 methylenedioxyphenyl) - 8 - acetoxymethyl- 8' hydroxymethyltetrahydrofuran, 7, 8 – trans - 8, 8' – trans - 7', 8' – cis -7- (5-methoxy-3,4 methylenedioxyphenyl) -7' -(4-hydroxy -3, 5-dimethoxyphenyl) -8, 8' diacetoxymethyltetrahydrofuran [30], sesamin [31], and isoswertisin [32]. Patuloside A (3- β -D-glucopyranosyloxy-1, 5, 6-trihydroxy-9H-xanthene-9-one) is a xanthone glycoside isolated from Peperomia pellucida. [1, 4, 33-40].

October - December 2011 RJPBCS Volume 2 Issue 4 Page No. 361



PHARMACOLOGICAL PROPERTY

Crude methanolic extracts of *Peperomia pellucida* has been reported as broad spectrum antimicrobial activity which was evaluated by the disk diffusion method. The fractions were found to be more active than the crude extracts [18]. Other studies document similar results for activity against numerous species, including Bacillus subtilis, Escherichia coli, Pseudomonas aeruginosa, and Staphylococcus aureus [41, 42]. Chloroform extracts from dried leaves of Peperomia pellucida, particularly the isolated compounds like apiol and pachypophyllin, has been reported a potent antifungal activity against Trichophyton mentagrophytes [35]. Antiprotozoal activity also reported in Peperomia pellucida plant. A whole plant extract inhibited growth of the chloroquine-resistant Plasmodium falciparum Indo strain by 95% in vitro at 100 mg/mL, and the rodent malaria Plasmodium vinckei petteri by 78% in vivo at 1,000 mg/kg. Anti-inflammatory and analgesic activity has been reported on aqueous extract of the aerial part of *Peperomia pellucida* plant. The activity was tested in rats and mice, respectively. Oral administration of 200 and 400 mg/kg of the aqueous extract exhibited an antiinflammatory activity whereas 400 mg/kg of the plant extract had the highest analgesic activity [11, 18, 19]. A neuropharmacological effect of *Peperomia pellucida* leaves has been reported in mice. Both petroleum ether and ethyl acetate fractions of ethanol extract of Peperomia pellucida leaves contain psychoactive substances which are CNS depressant in nature. In petroleum ether fraction was more active than ethyl acetate fraction and at the same doses, effects of petroleum ether fraction on the duration of diazepam-induced sleep and latency of the death caused by nikethamide toxicity was better [22]. The antipyretic Activity of Peperomia pellucida Leaves in Rabbit has been reported. Antipyretic effects of petroleum ether and ethyl acetate soluble fractions of ethanol extract of the leaves of Peperomia pellucida were intra peritoneal (i.p.) administration at a dose of 80 mg/kg body weight significantly reduced the elevated body temperature of rabbit [43]. Cytotoxicity was also observed in crude extracts from Peperomia pellucida against the cancer cell lines HL-60, MCF-7, and HeLa [44]. Peperomia pellucida constituents' flavonoids have been known to possess anti-oxidant, anti-neoplastic, anti-ulcer, anti-inflammatory and anti-microbial activities. This plant also reported at dosedependent increase in adverse effects in the major systems of the body such as integunrentarym, usculo-skeletal, nervous, respiratory, digestive and urogenital, covering the dose range from 6 g to 32 g per kg body weight of mouse along with a delayed appearance of adverse effects such as delayed time of death, delayed appearance of soft feces, and delayed recovery or no recovery from weight loss [45].

CONCLUSION

From the foregoing accounts it is evident that the plant *Peperomia pellucida* has been Ethno-medicinally used as a valuable therapeutic agent for a variety of diseases, as we have illustrated in this article. Moreover, numerous research works have proven its uses beyond the ethno-medicinal ones in experimental animals. Various compounds which were isolated from this plant may be responsible for its pharmacological activities. Being such a most useful and immense medicinal values that require more exploration in all the pharmaceutical aspect. So it needs further researches towards the development of safe and suitable medications. The road



ahead is to establish specific bioactive molecules, which might be responsible for these pharmacological actions.

ACKNOWLEDGEMENT

Department of Pharmacognosy, Rajiv Gandhi Institute of Pharmacy, Trikaripur, Kasaragod, Kerala- 10.

REFERENCES

- [1] Dos Santos PR, de Limas Moreira D, Guimaraes EF, Kaplan MA. Phytochem 2001; 58:547-551.
- [2] de Fatima Arrigoni-Blank M, Dmitrieva EG, Franzotti EM, Antoniolli AR, Andrade MR, Marchioro M. J Ethnopharmacol. 2004; 91: 215-218.
- [3] Arrigoni-Blank Mde F, Oliveira RL, Mendes SS, et al. BMC Pharmacol 2002; 2: 12-19.
- [4] Bayma JD, Arruda MS, Müller AH, Arruda AC, Canto WC. Phytochemistry 2000; 55:779-782.
- [5] Ghani A. Medicinal plants of Bangladesh. Bangladesh, Asiatic Society of Bangladesh, 1998, pp 77-78.
- [6] Flowers of India. http://www.flowersofindia.in/catalog/slides/Shiny%20Bush.html
- [7] United States Department of Agriculture. Natural Resources Conservation Service. http://plants.usda.gov/java/profile?symbol=PEPE5.
- [8] Wagner WL, Herbst DR, and Sohmer SH. Manual of the flowering plants of Hawai'i, (2 vols). Revised edition. Bernice P. Bishop Museum special publication. University of Hawai'i Press/Bishop Museum Press, Honolulu. 1999; pp 1034.
- [9] Santos PR, Moreira DL, Guimaraes EF, Kaplan MA. Phytochem 2001; 54: 547-551.
- [10] Mde FA, Oliveira RL, Mandes SS. BMC Pharmacol 2002; 2: 12-19.
- [11] Arrigoni-Blank MF, Dmitrieva EG, Franzotti EM, Antoiolli AR, Andrade MR, Marchioro M. J Ethnopharmacol 2004; 91: 215-218.
- [12] Smith, Albert C. Flora Vitiensis nova: a new flora of Fiji. National Tropical Botanical Garden, Lawai, Kauai, Hawaii. 1981; Volume- 2: pp 89.
- [13] Cheng YQ. Flora Reipublicae Popularis Sinicae; Science Press: Beijing, 1982; Vol. 20, p 77.
- [14] Whistler WA. Checklist of the weed flora of western Polynesia. Technical Paper No. 194, South Pacific Commission, Noumea, New Caledonia, 1988; pp 33.
- [15] Henty EE & Pritchard GH. Weeds of New Guinea and their control. 2nd Edition.
 Department of Forests, Division of Botany, Botany Bull. No.7, Lae, Papua New Guinea.
 1975; pp 133.
- [16] Craig WJ Health-promoting properties of common herbs. Am J Clin Nutr 1999; 70: 491S-499S.
- [17] Hua YX, Liu SF, Yang ZQ. Chinese Bencao; Shanghai Science & Technology Press: Shanghai, 1999; 3: 422.
- [18] Khan MR, Omoloso AD. Fitoterapia 2002; 73: 251-254.
- [19] Aziba PI, Adedeji A, Ekor M, Adeyemi O. Fitoterapia 2001; 72:57-58.
- [20] Bayma JC, Arruda MSP, Mu"ller AH, Arruda AC, Canto WC. Phytochem 2000; 55, 779-782



- [21] Muñoz V, Sauvain M, Bourdy G, et al. J Ethnopharmacol 2000; 71:123-131.
- [22] Khan A, Rahman M, Islam MS. DARU 2008; 16 (1):
- [23] Calimag MMP. Herb-Drug Interactions. Manila, Philippines, Manila Medical Society. 2007;
- [24] http://www.geocities.com/mmsi1902/herbal_aware.htm
- [25] Pansit-Pansitan (Peperomia pellucida Linn.) a.k.a. Ulasiman-Bato. Philippine Herbal Medicine. www.philippineherbalmedicine.org/pansit-pansitan.htm.
- [26] Ayurvedic Medicinal Plant. [http://ayurvedicmedicinalplants.com/plants/2180.html]
- [27] Mishra MP. Peperomia pellucida, an Amazing Wild Medicinal Herb. Ecosensorium. Org. November 28, 2010. http://www.ecosensorium.org/2010/11/peperomia-pellucida amazing-wild.html.
- [28] Egwuche RU, AA Odetola and OL Erukainure. Res J Phytochem 2011; 5: 48-53.
- [29] Chen CM, Jan FY, Chen MT, Lee TS. Heterocycles 1989, 29, 411-414.
- [30] Govindachari TR, Krishna Kumari GN, Partho PD. Phytochemistry 1998, 49, 2129-2131.
- [31] Wu JL, Li N, Hasegawa T, Sakai J, Kakuta S, Tang WX, Oka S, Kiuchi M, Ogura H, Kataoka T, et al. J Nat Prod 2005; 68: 1656-1660.
- [32] Hsieh TJ, Lu LH, Su CC. Biophys Chem 2005; 114: 13-20.
- [33] Webby RF, Markham KR. Phytochem 1994; 36: 1323-1326.
- [34] Xu S, Li N, Ning MM, Zhou CH, Yang QR, Wang MW. J Nat Prod. 2006; 69: 247-250.
- [35] Moreira DL, De Souza PO, Kaplan MA, Guimaraes EF. Acta Hortic 1999; 500: 65-69.
- [36] Ragasa CY, Dumato M, Rideout JA. ACGC Chem Res Commun 1998; 7: 54-61.
- [37] Agil M, Rahman FA, Ahmad MB. Sci Phys Sci 1994; 6: 141-143.
- [38] Agil M, Khan IZ, Ahmad MB. Sci Phys Sci 1993; 5: 213-215.
- [39] Manalo JB, Han BH, Han YN, Park MH, Anzaldo FE. Arch Pharm Res 1983; 6:133-136.
- [40] Oliveros-Belardo L. Perfum Essent Oil Rec 1967; 58: 359-363.
- [41] da Silva MH, Zoghbi MG, Andrade EH, Maia JG. Flavour Fragrance J 1999; 14:312-314.
- [42] Bojo AC, Albano-Garcia E, Pocsidio GN. Asia Life Sci 1994; 3: 35-44.
- [43] Akinnibosun HA, Akinnibosun FI, German BE. Sci world J 2008; 3 (4)
- [44] Alam Khan, Moizur Rahman, Shariful Islam. Turk J Biol 2008; 32: 37-41.
- [45] Chan-Bacab MJ, Peña-Rodriguez LM. Nat Prod Rep 2001; 18: 674-688.
- [46] SusieO Sio, Nelia P Cortes-Maramba S, I dro C Sia, Acute Oral Toxicity Of The Frbeze-Driedaqueous Extract Of Peperomiap ellucida (L) HBK (ulasimangb ato) in Mice; Acta Medica Philippina.