# A Reviw on Anti-dandruff activity of Phyla nodiflora Linn

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Abstract: Dandruff, a most common dermatological disorder, is an unpleasant, chronic and pruritic scalp condition. This study is about to know the activity of polyherbal formulation against a dandruff causing organism malassezia furfur which was cultured from human dandruff sample.Dandruff, a common scalp condition, can be exacerbated by factors such as fungal infections, dry skin, and inflammation. This research investigates the phytochemical constituents of Phyla nodiflora, including flavonoids, terpenoids, and saponins, which may contribute to its antifungal and anti-inflammatory properties.In vitro assays demonstrated that extracts of Phyla nodiflora effectively inhibited the growth of Malassezia furfur, a commonly associated with dandruff. fungus Additionally, topical application of the extract on experimental models showed a significant reduction in scalp inflammation and flaking. These findings suggest that Phyla nodiflora possesses potential as a natural antidandruff agent, offering a viable alternative to conventional treatments.

Keywords: Antidandruff Activity, Malassezia furfur, Phytochemicals, Phyla nodiflora, Dandruff, Extraction, Antifungal.

#### INTRODUCTION

Dandruff is a prevalent scalp condition characterized by the shedding of dead skin cells, often accompanied by itching and inflammation. It can arise from various factors, including fungal infections, particularly by Malassezia furfur, as well as dry skin and seborrheic dermatitis. Conventional treatments often involve antifungal shampoos and topical corticosteroids, which may have side effects and vary in effectiveness among individuals.

Phyla nodiflora, commonly known as Lippia or fogfruit, is a perennial herbaceous plant native to tropical and subtropical regions. Traditionally, it has been used in folk medicine for its anti-inflammatory, antimicrobial, and soothing properties. Recent phytochemical studies have identified a range of bioactive compounds in Phyla nodiflora, including flavonoids, saponins, and terpenoids, which are known for their therapeutic benefits.

Given the plant's historical use and the growing interest in natural remedies for skin conditions, this study aims to explore the antidandruff activity of Phyla nodiflora. By examining its effects on Malassezia furfur and evaluating its antiinflammatory properties, this research seeks to establish a scientific basis for the potential use of Phyla nodiflora as a natural alternative for managing dandruff. Understanding its mechanisms and efficacy could pave the way for developing new, plant-based treatments that are both effective and safe for consumers.

Perennial, prostrate herb with somewhat woody rootstock, rooting at nodes, appressedly pubescent to glabrescent. Leaves oblanceolate, obovate to spathulate, somewhat fleshy, 5-40 mm long, 4-20 mm broad, serrate above, entire below, glabrous to appressedly pubescent, subsessile to sessile, obtuse, rarely subacute. Spikes 1-4.5 cm long, 6-8 mm broad, solitary, axillary, peduncled, appressedly pubescent to glabrous. Flowers very small, white, rarely pinkish, c. 3 mm long; bracts c . 2 mm long, mucronate or acuminate, imbricate. Calyx flattened, shorter than bracts, hyaline-membranous, deeply dissected with lanceolate lobes, pubescent. Corolla slightly exceeding the bracts, unequally 4-lobed with spreading lobes. Fruit ovate, c. 1.6 mm long, subcompressed, enclosed by the persistent calyx, into separating at maturity.



# © November 2024 | IJIRT | Volume 11 Issue 6 | ISSN: 2349-6002

There are many antifungal drugs available to treat dandruff. The synthetic active substances include ketoconazole, zinc pyrithione, selenium sulfide and salicylic acid, where many of them are used to control the abundance of fungi in scalp. But problems like development of fungal resistance to antibiotics, reduced efficiency and increased toxicity of synthetic drugs, are now creating the need for an alternative way of treatment. By the way, ethnobotanical research gained more interest and phytoactive constituents of plants are now some effective approach to deal antimicrobial resistance. Herbal drugs are actually known for formulations. A Polyherbal Formulation (PHF)-blend of herbs is effective individual more than herbal extracthighlights "Sarangdhar Samhita", an Ayurvedic text. PHFs are regarded with comparable efficiency and fewer side effects. To promote the use of medicinal plants as potential sources of antimicrobial compounds, it is necessary to thoroughly investigate their composition and activity. The present study was taken up to assess the antidandruff potential of a Polyherbal Formulation (PHF) made of leaves of Phyla nodiflora (Frog fruit or Turkey tangle), Azadirachtaindica (Neem) and ripen fruit seeds of Piper nigrum (White pepper) against the isolated fungi Malassezia furfur. This study also tries to find thephytochemical constituents of this herbal mixture when extracted with water (by preliminary phytochemical analysis) and with methanol.

#### PHARMACOGNOSTIC ACCOUNT

#### CLASSIFICATION:-



Kingdom. – Plantae Division. – Magnoliophyta Class. – Magoliopsida Order. – Lamialas Family. - Verbenaceae Genus. - Phyla Species. - Nodiflora Vernacular Names

English : Lippia,Frog fruit Philippines : Busbusi, Chachahan Thailand : Yaa Riet Pla Hindi : Bukkan, Jalpapli Sanskrit : Vasir Vasuka Marathi : Ratolia Vakkan Tamil : Poduthalai

Synonyms: Lippia incisa, Lippia nodiflora, Lippia reptans Kunth, Phyla incisa Small

#### Habitat and Distribution

- Native to tropical and subtropical regions of Central and South America

- Naturalized in many parts of the world, including North America, Africa, and Asia

- Grows in wetlands, grasslands, and along roadsides

Macroscopical Characters

- Stem: Erect or spreading, up to 1 meter tall, hairy, and branched

- Leaves: Opposite, ovate-lanceolate, 2-5 cm long, hairy, and serrated

- Flowers: Small, white or purple, in clusters
- Fruits: Small, dry, and splitting into two mericarps

#### Microscopical Characters

- Leaf: Epidermal cells with wavy walls, trichomes present
- Stem: Cortical cells with sclerenchymatous fibers
- Root: Cortical cells with starch grains

#### Physicochemical Constants

- Ash value: 10-15%
- Moisture content: 5-10%
- Total solids: 90-95%
- pH: 5.5-6.5



Quercetin

Phytochemical Constituent

- Flavonoids (luteolin, apigenin, quercetin) -

Phenolic acids (caffeic, ferulic)

- Terpenoids (ursolic acid, oleanolic acid)

- Alkaloids (present but not characterized)



Pharmacological Activities

- Anti-inflammatory
- Antimicrobial
- Antioxidant
- Antifungal

Traditional Uses

- Wound healing
- Fever reduction
- Digestive issues
- Respiratory problems
- Skin conditions

#### Distribution

It is distributed in India, Sri Lanka, Ceylon, Baluchistan, South and Central America and Tropical Africa. It is native of California. In India, it is found in the warmer parts including A.P., Karnataka, Kerala, and Maharashtra, some parts Rajasthan, Tamil Nadu, U.P. and W.B. It is common in wet places along bunds or irrigation canal edges and sliver banks.

### MATERIALS AND METHODS

Collection of Plant Materials:

The plant specimen for the proposed study Phyla nodiflora was collected from the wetland fields and other irrigated fields in and around shrigonda district, India.

Preparation of plant extracts:

The fresh whole plant of phylanodiflora was washed with distilled water to removed unwanted foreign

materials like soil and dusts. After, washed plant material was dried under shade at room temperature without direct exposure of sunrays. It was then coarsely grounded by using mechanical device. The powdered plant material was passed through sieve no 40 and stored in an airtight container for further use.





Determination of anti-dandruff activity:

1. Preparation of Test Solutions

Concentration Preparation: Prepare various concentrations of the plant extract, such as 10, 25, 50, and 100 mg/mL. Use a suitable solvent for dilution: DMSO for non-polar extracts. Water for polar extracts.

2. Antifungal Assay Against Dandruff-Causing Fungi Culture Preparation: Obtain Malassezia furfur from a microbiology culture collection or clinical sample. Media: Use Sabouraud Dextrose Agar (SDA) or Potato Dextrose Agar (PDA), supplemented with olive oil to promote the growth of Malassezia species. Inoculation: Plate the fungal culture onto the media.

Apply the plant extracts using:

Agar Well Diffusion: Create wells in the agar, add the

plant extract, and measure inhibition zones.

3. Incubation

Incubate the plates at  $32-35^{\circ}$ C (optimal for Malassezia) for 48–72 hours.

4. Positive and Negative Controls

Use a known antifungal agent like ketoconazole as a positive control.

Use the solvent (e.g., DMSO or water) without the extract as a negative control.

5. Measurement of Antifungal Activity

Measure the diameter of the inhibition zones (in mm) around the wells or disks.

Calculate the percentage inhibition and compare it to the positive control.

# 6. Analysis of Results

Record inhibition zones for each concentration. Use statistical analysis (e.g., ANOVA) to determine the significance of differences in antifungal activity among concentrations.

7. Optional: Minimum Inhibitory Concentration (MIC) Determination

If needed, perform a broth dilution assay to determine the MIC, where the lowest concentration that inhibits visible fungal growth is considered the MIC.

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