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A Review on Extraction, Isolation and Separation Technique Studies of Curry Leaves

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Abstract: *Murraya koenigii, family Rutaceae, commonly known as Curry leaf plant is a highly valued plant for its medicinal value and characteristic aroma. The plant is a rich source of carbazole alkaloids. The petroleum ether, chloroform, ethyl acetate and ethanol extracts of roots of the plant were screened for phytochemical properties and antimicrobial activity for Staphylococcus aureus, Micrococcus luteus, Bacillus subtilis, Escherichia coli, Pseudomonas aeruginosa, Candida albicans and Aspergillus niger. Phytochemical screening showed the presence of carbohydrates, alkaloids, steroids and flavonoids in the root extracts of the plant. The study shows that all the extracts possess remarkable antibacterial activity. Additionally, petroleum ether and chloroform extracts also had antifungal activity.*

Keywords: Murraya koenigii, Rutaceae, phytochemical screening, antimicrobial activity

I. INTRODUCTION

Murraya koenigii (Linn,) Spreng. (Family-Rutaceae) commonly called Curry leaves in trade, occurs throughout India up to an altitude of 1500 metres (1). The leaves of the plant are used as a flavoring agent.Medicinally, these leaves found use in diarrhea, dysentery and to prevent vomiting. The Leaves and fruits are also a source of an essential oil which finds use as a fixative for heavy type of soap perfume. Leaves, root and bark are tonic, stomachic and carminative. Juice of roots provides relief from renal pain (8). Previous phytochemical investigations on this plant revealed the presence of carbazole alkaloids (3, 6, 10, 12, 13, 14, 15, 16) and coumarins (2). The present study is aimed at preliminary phytochemical screening of the root extracts of Murraya koenigii and evaluation of the same for potential antimicrobial activity**.**

Extraction Techniques

Extraction is a critical initial step in isolating bioactive compounds from Curry leaves Several techniques have been utilized, each with its advantages and limitations, depending on the target compounds and the desired yield.

1. Maceration: Maceration is one of the oldest and simplest extraction methods, involving soaking plant materials in solvents at ambient temperature. The process is straightforward and requires minimal equipment, making it accessible. However, maceration has drawbacks, such as long extraction times and the potential for lower yields. For Moringa oleifera, maceration is effective in extracting phenolic compounds, though it may require high solvent volumes and longer durations to achieve satisfactory yields.

Procedure of Maceration

The plant material is finely ground or crushed to increase the surface area for better solvent penetration.
The material is then submerged in a solvent such as water, ethanol, methanol, or a solvent mixture.
The solvent is kept in contact with the material for an extended period, typically ranging from a few hours to several days, depending on the type of material, solvent, and target compounds.
Stirring or occasional shaking may be applied to enhance the extraction process.
After maceration, the solvent containing the extracted compounds (the filtrate) is separated from the solid plant material by filtration or decantation.

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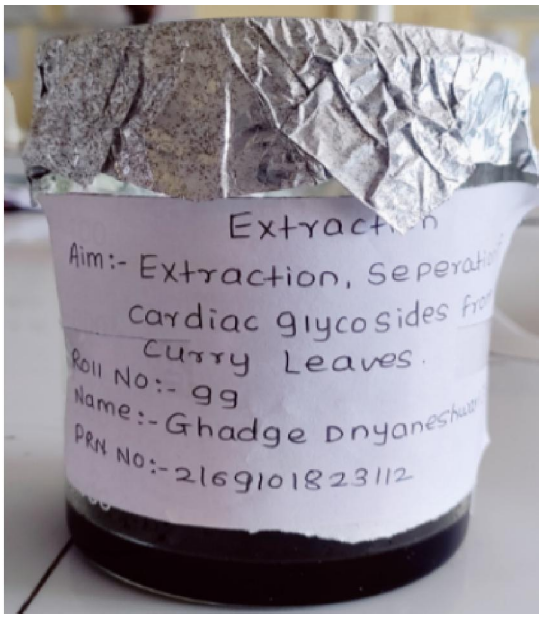
The solvent is often evaporated to obtain a concentrated extract, which can then be used for further studies or applications.



Maceration Process

Soxhlet Extraction: This method involves a continuous solvent reflux, where the solvent repeatedly passes through the plant material, enhancing extraction efficiency. Soxhlet extraction is widely used due to its high yield potential for heat-stable compounds. However, it requires long extraction times and large amounts of organic solvents, which may limit its environmental sustainability and cost-effectiveness. Soxhlet extraction has been effectively used to isolate various bioactive compounds from curry leaves including essential oils and other phenolic constituents.

Process:

The round-bottom flask containing the solvent is heated, causing the solvent to evaporate and travel upwards.

The vapor reaches the condenser, where it cools and condenses, then drips into the extractor chamber containing the plant material.

The solvent gradually fills the chamber, allowing the target compounds to dissolve into the solvent.

Once the chamber reaches a certain level, it siphons back down to the flask, carrying the dissolved compounds with it.

cycle repeats continuously, with fresh solvent contacting the plant material until the extraction process is complete.

Ultrasound-Assisted Extraction (UAE): UAE utilizes ultrasonic waves to disrupt cell walls, enhancing the release of bioactive compounds. The method is particularly advantageous due to its speed, low solvent requirement, and ability to extract heat-sensitive compounds, as it operates at lower temperatures. UAE has shown high efficacy in extracting antioxidants, phenolics, and essential oils from Curry leaves, making it a suitable choice for preserving delicate compounds while reducing extraction time and energy costs.

Procedure of Ultrasound-Assisted Extraction

Preparation: The plant material is usually ground or cut into smaller pieces to increase surface area.

Sonication Setup: The sample is placed in an extraction vessel with a suitable solvent, and an ultrasonic probe or bath generates ultrasonic waves.

Extraction Process: Ultrasonic waves are applied for a specified duration, typically ranging from a few minutes to an hour, depending on the plant material, solvent, and target compounds.

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Filtration and Concentration: After extraction, the solvent containing the dissolved compounds is filtered to separate the solid residue. The solvent may then be evaporated to yield a concentrated extract.

Microwave-Assisted Extraction (MAE): MAE employs microwave energy to heat solvents and plant materials, improving the efficiency of the extraction process. The method is known for its rapid processing times, reduced solvent usage, and high yields of bioactive compounds. MAE has been successfully applied in extracting Curry leaves bioactive compounds, such as glucosinolates and saponins, due to its ability to disrupt plant cell matrices and release intracellular compounds effectively. One limitation is the potential degradation of heat-sensitive components if the temperature is not carefully controlled.

Procedure of Microwave-Assisted Extraction

Preparation: The plant material is often dried and ground to increase surface area.
Microwave Setup: A microwave reactor or microwave-assisted extraction system is used, containing the plant material and solvent in an extraction vessel.
Extraction Process: The vessel is subjected to microwave irradiation, rapidly heating the solvent and plant matrix for a specified time (usually a few minutes).
Filtration and Concentration: After extraction, the solvent is filtered to separate the plant residue. The solvent may then be evaporated or further processed to concentrate the extract.

Supercritical Fluid Extraction (SFE): SFE uses supercritical fluids, typically carbon dioxide (CO_2), to extract non-polar compounds, offering selectivity, rapid extraction times, and minimal solvent residue. SFE is particularly effective for extracting lipophilic compounds such as essential oils and lipids Cymbopogon, and CO_2's non-toxic and non-flammable nature makes it an environmentally friendly choice. However, SFE requires specialized equipment and may have limited effectiveness for polar compounds unless co-solvents are used.

Procedure of Supercritical Fluid Extraction

Preparation: The plant material is dried and ground to increase the extraction surface area.
Supercritical Setup: The material is placed in an extraction chamber, and CO_2 is pumped under high pressure, becoming supercritical in state.
Extraction Process: Supercritical CO_2 passes through the plant matrix, dissolving target compounds. The CO_2 and extracted compounds then flow into a separation chamber, where CO_2 is depressurized back to its gaseous state.
Collection and Recovery: As CO_2 becomes a gas again, it releases the extracted compounds, which can then be collected. The CO_2 can be recycled for additional extractions, making the process environmentally efficient.

Hydrodistillation: Commonly used for extracting essential oils, hydrodistillation involves passing steam through the plant material to vaporize volatile compounds, which are then condensed and collected. Although efficient for isolating essential oils, hydrodistillation is limited in extracting non-volatile bioactive compounds. For Cymbopogon Citratus, hydrodistillation has been used to extract essential oils from seeds and leaves, though it may not be ideal for other bioactives.
Preparation: Plant material is often dried and sometimes chopped or ground to improve extraction efficiency.
Loading: The plant material is loaded into a distillation chamber with either water or directly exposed to steam, depending on the specific method.
Heating and Extraction: The chamber is heated, releasing essential oils and volatile compounds as steam.
Condensation and Separation: The vaporized essential oils and water pass through a condenser, turning into a liquid that collects in a separator. Since essential oils are typically immiscible with water, they form a separate layer that can be easily isolated.

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Isolation and Separation Techniques

Once the compounds are extracted, further isolation and separation are required to purify specific bioactive compounds. Techniques used for Moringa oleifera include chromatography, electrophoresis, and other advanced methods, each with varying degrees of specificity and resolution.

1. Chromatography: Chromatography techniques, including thin-layer chromatography (TLC), high-performance liquid chromatography (HPLC), and gas chromatography (GC), are widely used for separating and purifying bioactive compounds. HPLC is particularly effective in isolating Moringa compounds due to its high resolution and versatility in separating different types of compounds based on polarity and molecular weight. TLC is often used for initial screening and rapid qualitative analysis, while GC is suitable for volatile compounds such as essential oils. Chromatographic methods have been widely applied in isolating citral, geraniol, isogeranial and citronellol from Cymbopogon extracts.

2. Electrophoresis: Electrophoretic techniques, such as capillary electrophoresis (CE), offer high-resolution separation based on the charge-to-size ratio of molecules. Capillary electrophoresis is suitable for analyzing Cymbopogon smaller bioactive compounds and offers rapid analysis with minimal sample and solvent requirements. However, it may be less effective for larger, non-ionic compounds, limiting its application for certain Cymbopogon constituents.

3. Liquid-Liquid Extraction (LLE): LLE separates compounds based on their solubility in different solvents. It is a simple yet effective technique for fractionating Cymbopogon extracts into polar and non-polar components. LLE has been used as a preliminary separation step before further purification, particularly for isolating hydrophilic and lipophilic compounds in Cymbopogon extracts.**4. Preparative HPLC:** This technique is an advanced form of HPLC used to isolate large quantities of purified compounds for further study or application. Preparative HPLC has proven effective in isolating Cymbopogon Citratus bioactive constituents with high purity, though it requires sophisticated equipment and expertise.

II. CONCLUSION

Curry leaf extract contains alkaloids, flavonoids, saponins, polyphenols, and tannins as secondary metabolite chemicals. This makes the curry plant a potential candidate for traditional medicinal ingredients.

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