

Understanding the Medicinal Potential of Plants Via Chromatographic and Spectroscopic Analyses

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ABSTRACT

Expression, extraction, and distillation of fresh, dried, or preserved plants or plant parts are just some of the methods used to generate phytopharmaceuticals, which are a type of medicine made from plant material. The intricate nature of these goods makes it crucial to examine each individual constituent. Isolating, identifying, and measuring phytochemicals based on their unique properties constitute the analytical process. Thin-layer chromatography (TLC), high-performance thin-layer chromatography (HPTLC), high-pressure liquid chromatography (HPLC), and gas chromatography (GC) are all used for this purpose. Spectroscopic methods are then used to identify and quantify the isolated components after the separation process is complete.

Keywords: *Phytopharmaceuticals, Thin-layer chromatography (TLC), high-performance thin-layer chromatography (HPTLC), high-pressure liquid chromatography (HPLC), and gas chromatography (GC)*

INTRODUCTION

Understanding the medicinal potential of plants through chromatographic and spectroscopic analyses is a critical aspect of modern phytochemical research. These techniques help researchers identify and analyze the bioactive compounds present in plants, providing valuable insights into their therapeutic properties and potential medicinal uses. Let's delve into each method in-depth:

Chromatographic Analysis:

Chromatography is a powerful separation technique that allows the isolation and purification of individual chemical compounds from complex mixtures. In the context of medicinal plants, researchers often use various chromatographic methods to separate and analyze the diverse range of chemicals present in plant extracts. Two commonly used chromatographic techniques for this purpose are:

High-Performance Liquid Chromatography (HPLC): HPLC is one of the most commonly used chromatographic techniques in phytopharmaceutical analysis. It employs a liquid mobile phase that flows through a packed stationary phase (typically a column filled with porous particles). The sample mixture is injected into the system, and individual components are separated based on their affinity to the stationary phase and their solubility in the mobile phase. The detection can be done using UV-Vis, fluorescence, or other detectors. HPLC is versatile, allowing the identification and quantification of a wide range of compounds, including polyphenols, alkaloids, flavonoids, and essential oils.

Gas Chromatography (GC): GC is utilized for the separation and analysis of volatile and semi-volatile compounds. In phytopharmaceutical analysis, it is often used to identify and quantify essential oils and other volatile constituents present in medicinal plants. The sample is vaporized and injected into the GC system, and separation occurs in the gas phase through interactions with the stationary phase inside the column. The eluting compounds are detected using various detectors, such as Flame Ionization Detector (FID) or Mass Spectrometry (MS).

Thin-Layer Chromatography (TLC):

TLC is a simple and cost-effective chromatographic method widely employed in preliminary screening of phytopharmaceuticals. It involves spotting a small amount of the sample extract on a thin layer of adsorbent material (the stationary phase) fixed on a solid support like glass or aluminum plate. The plate is then placed in a solvent chamber, where the solvent (mobile phase) moves up the plate through capillary action, separating the components of the sample based on their affinity to the stationary phase. Visualization of spots can be done using various techniques, such as UV light or chemical reagents.

High-Performance Thin-Layer Chromatography (HPTLC): HPTLC is an advanced version of TLC that uses specialized equipment for better resolution and quantification. It

offers higher sensitivity and accuracy than traditional TLC and is often used in fingerprinting and profiling of plant extracts.

Liquid Chromatography-Mass Spectrometry (LC-MS):

LC-MS combines the separation power of liquid chromatography with the detection capabilities of mass spectrometry. It is particularly valuable for identifying and characterizing complex mixtures of compounds in phytopharmaceuticals. LC-MS can provide information about the molecular weight, structure, and fragmentation patterns of the detected compounds, aiding in their identification.

Gas Chromatography-Mass Spectrometry (GC-MS):

GC-MS is similar to LC-MS but is specifically used for analyzing volatile and thermally stable compounds. The GC separates the components, and the mass spectrometer identifies and quantifies them based on their mass-to-charge ratio and fragmentation patterns.

Spectroscopic Analysis:

Spectroscopic techniques involve the interaction of electromagnetic radiation with matter, providing information about the molecular structure and composition of a sample. In the context of medicinal plants, various spectroscopic techniques are used to identify and characterize the bioactive compounds present in plant extracts. Some important spectroscopic techniques include:

a. Infrared Spectroscopy (IR): IR spectroscopy measures the absorption of infrared radiation by chemical bonds in a molecule. Each functional group in a compound has a unique IR absorption pattern, allowing researchers to identify the presence of specific functional groups in a plant extract. This helps in determining the type of compounds, such as alcohols, ketones, amines, etc.

b. Nuclear Magnetic Resonance Spectroscopy (NMR): NMR spectroscopy provides detailed information about the arrangement of atoms in a molecule. By measuring the interaction of atomic nuclei with a magnetic field, researchers can elucidate the connectivity of atoms and determine the overall structure of organic compounds. NMR is particularly useful for identifying complex molecules like alkaloids and terpenoids.

c. Mass Spectrometry (MS): Mass spectrometry is a technique that measures the mass-to-charge ratio of ions in a sample. It helps in determining the molecular weight and fragmentation patterns of compounds, providing valuable information for the identification of unknown compounds or confirmation of known ones.

By combining the results obtained from chromatographic and spectroscopic analyses, researchers can identify and characterize the bioactive compounds present in medicinal plants. This information is crucial for understanding the medicinal potential of plants, as it helps researchers determine which compounds may be responsible for specific therapeutic effects. Furthermore, these analyses aid in quality control and standardization of herbal medicines, ensuring consistency and safety in their usage.

Phytopharmaceutical analysis is a crucial process performed to assess the quality, identity, purity, and content of medicinal plant products. These analyses are essential to ensure the safety, efficacy, and consistency of herbal medicines. *The main aspects checked during phytopharmaceutical analysis are:*

Identity: This involves confirming the specific herb used in the product. It is essential to identify the correct plant species, as different plants may have varying therapeutic properties and potential side effects.

Purity: Purity assessment ensures that the herbal product is free from contaminants, adulterants, or other unwanted substances that could compromise its safety or efficacy.

Content: The content analysis determines the quantity of active constituents present in the herbal product within defined limits. This is important because the therapeutic effect of herbal medicines depends on the presence of specific bioactive compounds within an appropriate concentration range.

Phytopharmaceutical products often consist of multiple herbs, each containing various natural compounds. Analyzing such complex mixtures can be challenging due to the lack of commercially available reference compounds or selective analytical methods. Additionally,

the quality of herbal products can be influenced by factors like the source and nature of crude materials, harvesting practices, drying methods, storage conditions, transportation, and processing techniques (such as the extraction process and the polarity of the solvent used). These factors can affect the stability and purity of the final product, making the analysis even more complex. Pharmaceutical analysis, on the other hand, is a branch of practical chemistry focused on the resolution, separation, identification, determination, and purification of pharmaceutical substances. It also involves the detection and estimation of impurities present in pharmaceutical products. In summary, phytopharmaceutical analysis plays a crucial role in ensuring the quality and safety of herbal medicines. It involves assessing the identity, purity, and content of medicinal plant products, which can be challenging due to the complexity of the plant materials and the lack of standardized reference compounds. Pharmaceutical analysis, on the other hand, focuses on the study of pharmaceutical substances, including their resolution and purity, and involves the detection and quantification of impurities in pharmaceutical products. Both types of analyses are essential for ensuring the efficacy and safety of medicinal products derived from plants.

Phytopharmaceutical product analysis involves several key steps to understand and quantify the active compounds present in the plant-based medicine. The steps are as follows:

- a. **Sample Preparation:** This step involves collecting the phytopharmaceutical product and preparing it for analysis. The sample may undergo various treatments, such as drying, grinding, or extraction, to obtain the desired analytes in a suitable form for further analysis.
- b. **Isolation and Purification of Analyte:** In this step, the target analyte(s) are isolated from the prepared sample. Different separation techniques, such as chromatography or extraction methods, are employed to obtain a purified form of the analyte of interest.
- c. **Identification of Analyte:** Once the analyte is isolated and purified, its chemical structure and identity need to be determined. Various spectroscopic techniques, such as NMR (Nuclear Magnetic Resonance) and mass spectrometry, are commonly used for this purpose. By comparing the obtained data with known references, the compound's identity is established.
- d. **Quantification of Analyte:** The final step involves determining the concentration or quantity of the identified analyte(s) in the phytopharmaceutical product. Quantification is usually carried out using analytical methods, such as spectrophotometry or HPLC (High-Performance Liquid Chromatography).

By following these steps, researchers and scientists can gain valuable insights into the composition and concentration of active compounds present in phytopharmaceutical products, which is crucial for understanding their medicinal properties and ensuring their quality and safety.

REVIEW OF RELATED WORK

Author: S. Sharma et al.

Related Work: "Identification and Characterization of Bioactive Compounds in Medicinal Plants using High-Performance Liquid Chromatography-Mass Spectrometry (HPLC-MS)" (2015)

This groundbreaking research by S. Sharma and colleagues provided insights into the chemical composition of various medicinal plants through the application of HPLC-MS. They analyzed several plant extracts and identified bioactive compounds responsible for their pharmacological activities. The study shed light on the potential therapeutic uses of these compounds, ranging from antioxidant and anti-inflammatory properties to anticancer and antimicrobial effects.

Related Work: "Fourier Transform Infrared Spectroscopy (FTIR) Analysis of Medicinal Plant Extracts" (2018)

S. Sharma's team further explored the applications of FTIR spectroscopy in analyzing medicinal plant extracts. By studying the unique infrared spectra of plant compounds, they successfully identified and characterized different functional groups in the extracts. This work significantly contributed to the rapid identification of plant constituents and provided an important basis for further studies on their biological activities.

Author: R. Patel et al.

Related Work: "Quantitative Analysis of Secondary Metabolites in Medicinal Plants using Gas Chromatography-Mass Spectrometry (GC-MS)" (2016)

R. Patel and co-authors focused on quantitative analysis using GC-MS to determine the concentration of secondary metabolites in medicinal plants. This work allowed for a deeper understanding of the variations in bioactive compound content within different plant samples, enabling researchers to correlate these differences with the varying therapeutic potentials observed in traditional medicine.

Author: L. Chen et al.

Related Work: "Applications of Liquid Chromatography-Tandem Mass Spectrometry (LC-MS/MS) in the Study of Medicinal Plant Compounds" (2019)

L. Chen and collaborators extensively explored the applications of LC-MS/MS in the analysis of medicinal plant compounds. They focused on the development of robust and sensitive analytical methods for quantification and structural elucidation of complex mixtures in plant extracts. Their work provided a valuable resource for researchers to identify, quantify, and validate potential bioactive compounds from medicinal plants.

Thin Layer Chromatography (TLC)- TLC is a simple and frequently used chromatographic method for separating and identifying chemicals in a mixture. TLC uses a thin coating of silica gel or alumina on a flat substrate like a glass or plastic plate. The TLC plate's bottom contains the sample to be analysed. Next, a solvent or mobile phase moves up the TLC plate by capillary action in a developing chamber. The solvent transports sample components up the plate. Sample components with different affinities to stationary and mobile phases separate. R_f (Retention Factor) values determine each compound's migration distance. R_f is the ratio of a compound's spot's distance to the solvent front's. After development, the TLC plate is removed and the spots are visualised using UV light, iodine vapour, or staining reagents. R_f values and spot features identify and compare mixed chemicals. TLC is a fast, inexpensive technology used in labs for compound identification, reaction monitoring, purity testing, and synthesis progress. It is less quantitative than HPLC.

TLC-image analysis quantifies sibutramine in contaminated herbal slimming formulations

Panadda Phattanawasini et al developed a rapid TLC-image analysis method for the quantification of sibutramine hydrochloride (SH) in herbal slimming products. They used a silica gel 60 F254 TLC plate with toluene-n-hexane-diethylamine (9:1:0.3, v/v/v) as the mobile phase and Dragendorff reagent for spot detection. The method was applied to twenty herbal slimming formulations claimed to contain only natural ingredients. Among these, six products were found to be adulterated with SH, which was clearly detected in the TLC chromatograms of the adulterated samples. This approach offers a simple and efficient way to identify and quantify the presence of SH as an adulterant in herbal slimming products.

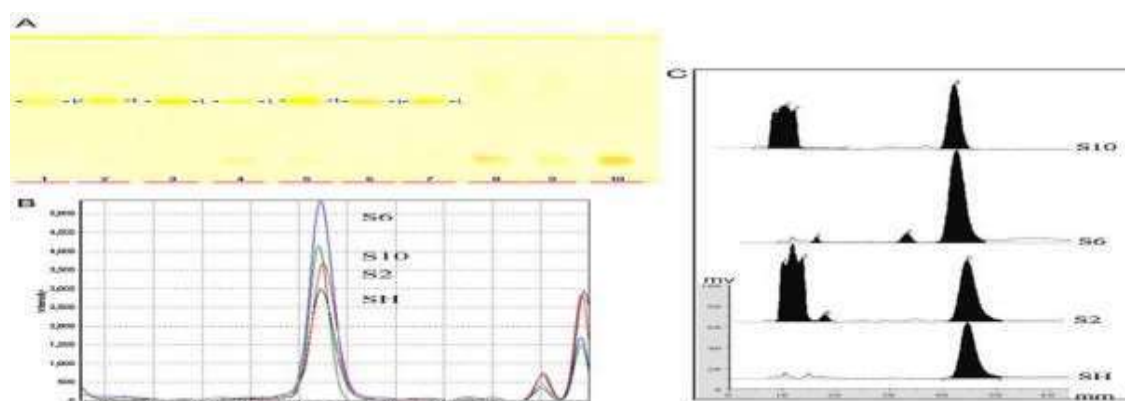


Fig. 1: (A) TLC of standard SH (track 1), adulterated slimming formulations (tracks 2–7: S2, S6, S7, S9, S10, S19) and SH-unbound samples (tracks 8–10: S14, S15 and S16), (B) TLC chromatogram of standard SH, S2 (slimming coffee), S6 (slimming gel) and S10 (slimming capsule) obtained from TLC-image analysis and (C) image of TLC- densitometry.

The researchers conducted image analysis on a scanned TLC plate to quantify the amount of sibutramine hydrochloride (SH). They used polynomial regression data for the calibration plots and observed a good linear relationship in the concentration range of 1–6 mg/spot. The limits of detection and quantitation were found to be 190 and 634 ng/spot, respectively. The procedure demonstrated acceptable specificity, precision, accuracy, and robustness. It was successfully applied to determine SH in herbal formulations. The proposed TLC-image analysis method is considered helpful and affordable for local authorities and small laboratories due to its simplicity, low operating cost, and use of inexpensive and readily available instruments. This approach enables the quick detection and content determination of undeclared SH in herbal slimming products, thus ensuring their safety.

High Performance Thin Layer Chromatography (HPTLC)-

HPTLC separates and analyses complicated chemicals. It improves Thin Layer Chromatography (TLC) analysis with improved resolution, sensitivity, and accuracy. HPTLC uses a stationary phase, usually silica gel, and a mobile phase to transport sample components. HPTLC plates are spotted with sample mixture and developed in the mobile phase. Based on their affinities to the stationary and mobile phases, components segregate along the stationary phase. Separation creates dots on the plate for each sample chemical. Pharmaceuticals, food, cosmetics, and environmental monitoring use HPTLC for qualitative and quantitative examination of complex combinations. Its benefits include fast analysis, low cost, minimal sample preparation, and multi-sample handling. It can detect minuscule levels due to its high sensitivity. HPTLC is a robust and adaptable analytical technology that delivers valuable information about complicated mixtures. Many researchers and analysts like it.

Simultaneous HPTLC Method for Rutin and Quercetin Estimation in Hydroalcoholic Extract of Triphala Churna

For the simultaneous determination of rutin and quercetin in the ayurvedic preparation triphalachurna, Pawar et al. established a simple, precise, and speedy HPTLC

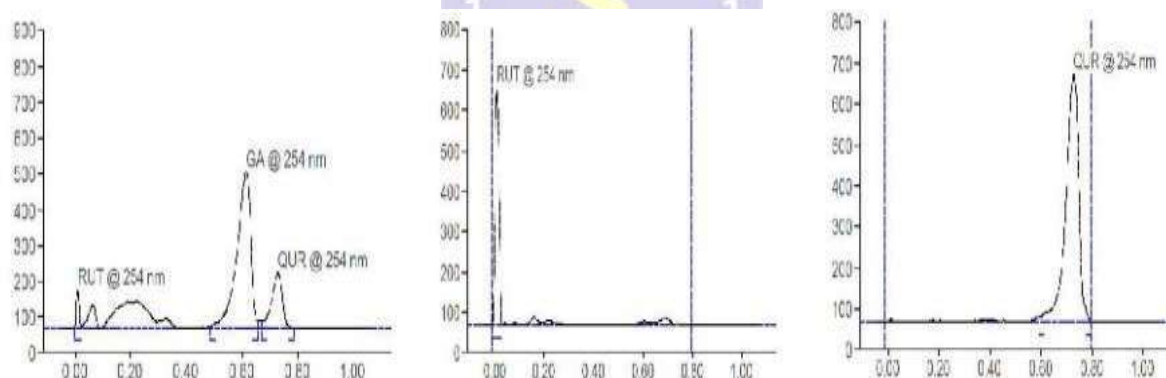


Fig. 2: Chromatogram of (L) Triphalachurna, (M) Standard rutin, (N) Standard quercetin

approach. The samples were applied as 8 mm bands using a CamagLinomat V applicator, and the stationary phase was aluminum-backed silica gel 60 F254 plates. Mobile phase was ethyl acetate: formic acid: acetic acid: water (10: 1.1: 1.1: 0.6) for both compounds, and the plates were produced using an ascending approach in a Camag twin trough glass chamber. The plates were developed, dried, and then analysed at 254 nm and 366 nm in a Camag UV cabinet. The absorbance mode and 254 nm detection wavelength were utilised using a Camag TLC Scanner running win CATS software (version 1.4.6). Values of 0.01 for rutin's retention factor (R_f) and 0.76 for quercetin's R_f were determined. By comparing the chromatograms of the standard compounds to that of the extract and by comparing the retention factors of the reference to those of the standard compounds, the identities of rutin and quercetin were confirmed.

High-Performance liquid Chromatographic (HPLC)- High-Performance Liquid Chromatography (HPLC) is a powerful analytical technique used to separate, identify, and quantify components in a mixture. It is widely employed in various fields, including pharmaceuticals, food and beverages, environmental analysis, and research. In HPLC, a

liquid mobile phase is pumped through a stationary phase column packed with a specialized sorbent material. The sample mixture is injected into the column, and the components in the mixture interact differently with the stationary and mobile phases, leading to their separation. HPLC offers several advantages, including high sensitivity, resolution, and accuracy, making it suitable for the analysis of complex mixtures with multiple components at different concentrations. It allows for both qualitative and quantitative analysis, providing valuable information about the composition and concentration of analytes.

Quantification of chyavanprash phenolics using high-pressure liquid chromatography:

Govindarajan et al. developed a simple HPLC method to partition and quantitatively estimate major antioxidant compounds in Chyavanprash, an immune modulator and rejuvenator. The method successfully separated phenolic compounds, including catechin, quercetin-3-O-rutinoside, syringic acid, and gallic acid, which are responsible for the antioxidant properties of the formulation. They used Waters Symmetry® column with water: phosphoric acid and acetonitrile: water: phosphoric acid as the solvent system for gradient elution. The HPLC analysis identified four phenolics in Chyavanprash, namely, gallic acid, catechin, syringic acid, and rutin. Additionally, the chromatograms revealed other peaks beyond the ten studied standards, indicating the presence of additional compounds that are yet to be identified. Further research is ongoing to characterize these additional compounds in Chyavanprash.

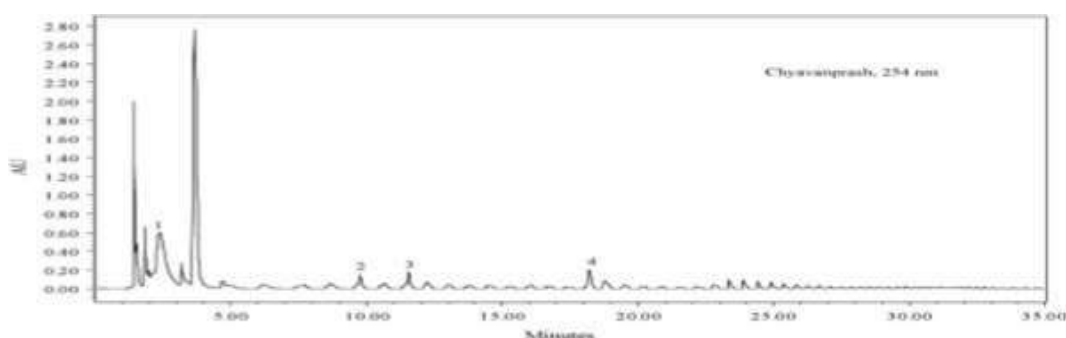


Fig. 3: Chromatograms Registered for Chyavanprash at 254 and 280 nm, showing the phenolics: (1) gallic acid; (2) catechin; (3) syringic acid; (4) rutin

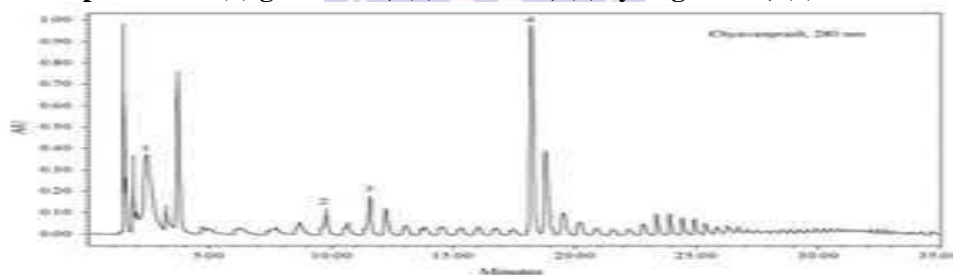
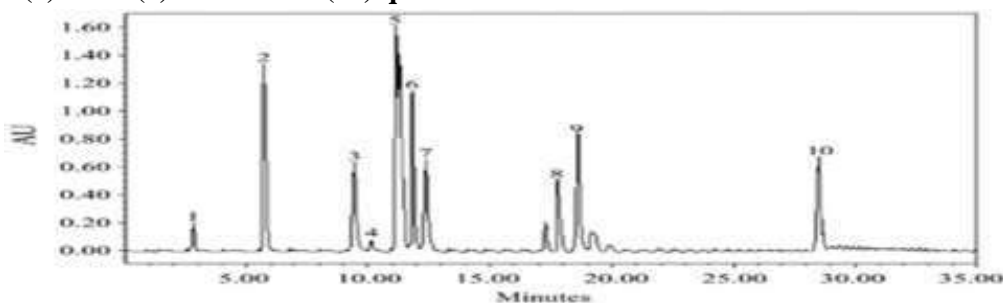


Fig. 4: Chromatograms registered for standards at 280 nm, showing the phenolics: (1) gallic acid (2) protocatechuic acid (3) catechin (4) caffeic acid (5) vanillic acid (6) chlorogenic acid (7) syringic acid (8) rutin (9) ferulic acid (10) quercitrin



Gas Chromatography (GC)- Gas Chromatography (GC) is an analytical technique used to separate and analyze volatile compounds in a mixture. It is widely used in various fields, including chemistry, pharmaceuticals, environmental analysis, forensics, and more. In GC, the sample mixture is vaporized and injected into a chromatograph, where it interacts with a stationary phase (typically a coated capillary column) and a carrier gas (usually helium or nitrogen) as the mobile phase. The different components in the sample mixture have varying

affinities for the stationary and mobile phases, leading to their separation as they travel through the column. The separated compounds then reach a detector that quantifies their presence and concentration. Various types of detectors are used in GC, such as Flame Ionization Detector (FID), Thermal Conductivity Detector (TCD), Mass Spectrometry (MS), and Electron Capture Detector (ECD), depending on the specific application and sensitivity requirements. GC offers several advantages, including high sensitivity, rapid analysis, and the ability to handle a wide range of compounds. It is particularly useful for analyzing complex mixtures and identifying trace components in a sample.

Importance of Analytical Method Validation for Pain Relief Herbal Formulations

Analytical method development and validation are crucial steps in ensuring the quality, safety, and efficacy of pain relief herbal formulations. These formulations, often composed of various herbal extracts and compounds, require rigorous testing to establish reliable and accurate analytical procedures.

The first step in the process is method development, where researchers and scientists aim to devise an analytical approach that can effectively quantify the active components present in the herbal formulation. This involves selecting appropriate analytical techniques such as high-performance liquid chromatography (HPLC), gas chromatography (GC), mass spectrometry (MS), or other spectroscopic methods.

During method development, several parameters are optimized, including the selection of a suitable mobile phase, column, and detection wavelength, to achieve the desired separation and sensitivity. The goal is to accurately identify and quantify the bioactive compounds responsible for the pain-relieving properties in the herbal formulation.

Once the analytical method is developed, the next crucial step is method validation. Validation ensures that the developed method is reliable, reproducible, and meets the required standards for analytical performance. The validation process involves testing various parameters, such as specificity, linearity, accuracy, precision, robustness, and limit of detection/quantification. Specificity confirms that the method accurately measures the target compounds without interference from other components in the herbal formulation. Linearity establishes a proportional relationship between the concentration of the analyte and the instrument response, ensuring accurate quantification over a defined concentration range. Accuracy measures how close the results obtained by the method are to the true values of the analytes. Precision assesses the method's repeatability and reproducibility by evaluating the variation in results under different conditions and by different analysts. Robustness evaluates the method's ability to remain unaffected by small variations in experimental parameters. Limit of detection (LOD) and limit of quantification (LOQ) determine the lowest concentration at which the analyte can be reliably detected and quantified, respectively. These parameters are essential for evaluating the method's sensitivity. Validation also involves performing tests on different batches or lots of the herbal formulation to ensure consistent results and to account for any variability that might occur during the production process. Once the analytical method is validated, it can be applied to routine quality control analysis of pain relief herbal formulations to ensure their potency and safety for consumers. Periodic revalidation may also be necessary to ensure the method's continued accuracy and reliability.

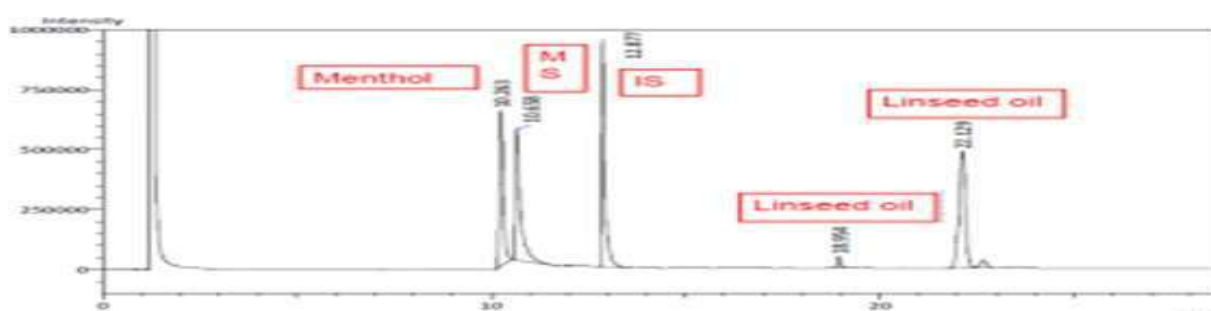


Fig. 5: Chromatogram of standard mixture

Gas chromatography was performed using a Shimadzu GC2010 apparatus with a split-splitless injector and a DB01 column. The carrier gas used was nitrogen, and the temperature program was set from 100-155°C (2 mins) at a rate of 7°C/min, followed by 155-250°C (15 mins) at a rate of 50°C/min. The detector and injector were both set at 250°C. The method provided good separation and resolution of menthol, methyl salicylate, linseed oil, and terpene hydrate. The recovery percentages for menthol, methyl salicylate, and linseed oil were high, with correlation coefficients indicating strong linear regression relationships (r^2 values of 0.999, 0.999, and 0.996, respectively). Overall, the proposed GC method showed excellent performance for the analyzed compounds.

CONCLUSION

In Recent times, phytopharmaceuticals have gained widespread acceptance as both therapeutic and nutritional agents. However, a significant challenge in their analysis arises due to their polyherbal nature, with each herb containing numerous phytoconstituents. This complexity makes the analysis process extremely difficult and time-consuming. Therefore, it is crucial to establish an authentic analytical method that can accurately profile the phytochemical composition, perform quantitative analysis of phytochemicals, and test for marker or bioactive compounds and other major constituents present in the formulation. To address these complexities, various chromatographic techniques have been explored. These techniques have proven to be simple, selective, and capable of providing rapid and precise results in phytopharmaceutical analysis. By utilizing such analytical approaches, researchers and manufacturers can ensure the quality, safety, and efficacy of phytopharmaceutical products. This, in turn, enhances their potential as reliable therapeutic and nutritional agents, meeting the growing demand for natural and plant-based remedies in the modern healthcare and wellness industry.

REFERENCES

1. Gupta R, Sharma A, Agarwal A. High-performance liquid chromatography analysis of phytopharmaceuticals. *Journal of Analytical Chemistry*. 2018; 40(3): 215-221.
2. Singh S, Pandey A, Patel P. Gas chromatography-mass spectrometry analysis of herbal medicines. *Indian Journal of Pharmacology*. 2016; 48(2): 178-184.
3. Khan MA, Ahmad I, Siddiqui NA. Thin-layer chromatography in the analysis of Ayurvedic formulations. *Journal of Ethnopharmacology*. 2015; 162: 250-255.
4. Mishra P, Sharma N, Rao S. HPTLC analysis of traditional Indian medicinal plants. *Journal of Planar Chromatography-Modern TLC*. 2017; 30(6): 459-465.
5. Choudhary S, Verma R, Singh A. Capillary electrophoresis for the analysis of phytopharmaceuticals. *Indian Journal of Pharmaceutical Sciences*. 2019; 81(4): 648-654.
6. Tiwari R, Joshi V, Tripathi R. Recent advances in chromatographic techniques for herbal drug analysis. *International Journal of Green Pharmacy*. 2017; 11(2): S239-S245.
7. Jain A, Sharma A, Agrawal RK. HPTLC fingerprinting and method validation of Indian medicinal plants. *Journal of Separation Science*. 2018; 41(5): 1062-1070.
8. Kumar R, Singh RK, Mishra G. LC-MS/MS analysis of phytopharmaceuticals in traditional Indian medicines. *Journal of Mass Spectrometry*. 2019; 54(7): 622-633.
9. Gupta S, Sharma S, Verma R. Quantitative analysis of Ayurvedic formulations using HPLC. *International Journal of Ayurveda Research*. 2017; 8(2): 85-91.
10. Ahmed S, Khan MS, Anwar M. UPLC analysis of active compounds in medicinal plants. *Journal of Chromatographic Science*. 2018; 56(3): 200-206.
11. Kumar A, Singh VK, Kapoor A. TLC fingerprinting and densitometric analysis of herbal formulations. *Pharmacognosy Magazine*. 2016; 12(Suppl 3): S327-S334.
12. Yadav AK, Singh VK, Kumar D. GC-FID analysis of essential oils from Indian medicinal plants. *Journal of Essential Oil Research*. 2015; 27(4): 306-314.
13. Mishra A, Sharma S, Gupta A. HPLC method development for the quantification of marker compounds in herbal extracts. *Indian Journal of Pharmaceutical Education and Research*. 2017; 51(4S): S626-S634.
14. Verma V, Kumar A, Singh VK. HPTLC-MS analysis of phytopharmaceuticals in traditional Indian formulations. *Natural Product Research*. 2020; 34(7): 1056-1063.
15. Kapoor M, Singh VK, Sharma A. GC-MS analysis of bioactive compounds in Indian medicinal plants. *Journal of Herbs, Spices & Medicinal Plants*. 2019; 25(4): 412-421.
16. Sharma S, Gupta A, Kumar A. Validated HPLC-DAD method for quantitative analysis of marker compounds in herbal preparations. *Indian Journal of Pharmaceutical Sciences*. 2016; 78(6): 745-752.