

Role of Chromatography in Evaluation of Herbal Drugs: A Short Review

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ABSTRACT

Herbal medicines are conventional as significant remedial agents for the healing of numerous conditions.

The development of genuine logical styles which can consistently outline the photochemical composition, as well as quantitative analyses of marker/bioactive composites and other foremost ingredients, is a major dispute to scientists. Pharmacognostical analysis of medicinal sauces remains grueling issues for logical druggists, as sauces are a complicated system of fusions. Analytical separation ways for illustration high performance liquid chromatography (HPLC) gas chromatography (GC) and group spectrometry (MS), High Performance Thin Layer Chromatography (HPTLC) etc. among the most popular styles of preference used for quality control of raw material and finished herbal product.

Key words– Herbal drug, Chromatography, Analysis.

I. INTRODUCTION

Chromatography represents the most protean partition fashion and readily available. Plant accoutrements are alienated and purified by using colorful chromatographic ways. Herbal drug is a difficult system of fusions. Therefore, the styles of preference for identification of 'botanical medicine' are substantially proposed to gain a characteristic point of a specific factory that signifies the presence of an exacting quality defining chemical ingredients. For similar purposes, chromatographic ways similar as high performance liquid chromatography (HPLC), gas chromatography (GC), gas chromatography – mass spectrometry

(GC- MS) and thin sub caste chromatography (TLC) were used extensively as reported in multitudinous publications¹.

Thin Layer Chromatography (TLC)

Thin layer chromatography is simply known as TLC. It is one of the most popular and simple chromatographic technique used of separation of compounds. In the phytochemical evaluation of herbal drugs, TLC is being employed extensively for the following reasons:

1. it enables rapid analysis of herbal extracts with minimum sample clean-up requirement,
2. it provides qualitative and semi quantitative information of the resolved compounds.
3. It enables the quantification of chemical constituents. Fingerprinting using HPLC and GLC is also carried out in specific cases.

In TLC characteristic, the data that can be recorded using a high- performance TLC (HPTLC) scanner includes the chromatogram, deceleration factor (R_f) values, the shade of the alienated bands, their immersion gamut's, λ maximum and shoulder curve/ s of all the determined bands. All of these, mutually with the biographies on derivatization with dissimilar reagents, represent the TLC point profile of the sample. The information so generated has an implicit operation in the recognition of an genuine medicine, in banning the pollutants and in maintaining the excellence and thickness of the medicine². TLC was the ordinary system of selection for herbal investigation before necessary chromatography styles like GC and HPLC were established. Indeed currently, TLC is still constantly used for the analysis of herbal drugs since colorful pharmacopoeias similar as American Herbal Pharmacopoeia (AHP)³, Chinese medicine studies and examination,

Pharmacopoeia of the People's Republic of Chianti. Still use TLC to give first attribute fingerprints of herbs⁴. Rather, TLC is used as an easier system of original webbing with a semi quantitative assessment together with other chromatographic ways. As there's fairly lower transform in the simple TLC partition of herbal drugs than with necessary chromatography, only a short review is given then, and for farther details about TLC the compendiums could consult references⁵⁻⁶.

High Performance Thin Layer Chromatography (HPTLC)

HPTLC fashion is extensively engaged in pharmaceutical assiduity in procedure development, recognition and discovery of pollutants in herbal product and helps in detection of fungicide content, mycotoxins and in excellence control of sauces and health Food.



Fig -1 High Performance Thin Layer Chromatography (HPTLC)

It has been well reported that several samples can be run contemporaneously by use of a low volume of mobile phase than in HPLC.

It has also been reported that mobile phases of pH 8 and over can be used for HPTLC.

Another benefit of HPTLC is the continual discovery (scanning) of the chromatogram with the same or dissimilar situation. Accordingly, HPTLC

has been delved for contemporaneous assay of several factors in amulet-component expression. With this fashion, authentication of colorful genus of factory possible, as well as the assessment of stability and thickness of their medications from different manufactures. Colorful workers have developed HPTLC system for phytoconstituents in crude medicines or

herbal phrasing similar as bergenin, catechine and gallic acid in *Bergenia cilliata* and *Bergenia lingulata*⁷.

High Performance Liquid Chromatography (HPLC)

Over the once decades, HPLC has entered the most expansive operation in the investigation of herbal drugs. Reversed-phase (RP) columns may be the most admired columns used in the logical partition of herbal drugs. Preliminary and logical HPLC are extensively used in pharmaceutical assiduity for segregating and sanctification of herbal composites. There are principally two types of preliminary HPLC: low pressure HPLC (generally under 5 bars) and high pressure HPLC (pressure > 20 bar).

The significant parameters to be considered are decision, perceptivity and fast investigation time in logical HPLC whereas both the degree of solute chastity as well as

the quantum of emulsion that can be produced per unit time i.e. outturn or recovery in preliminary HPLC.

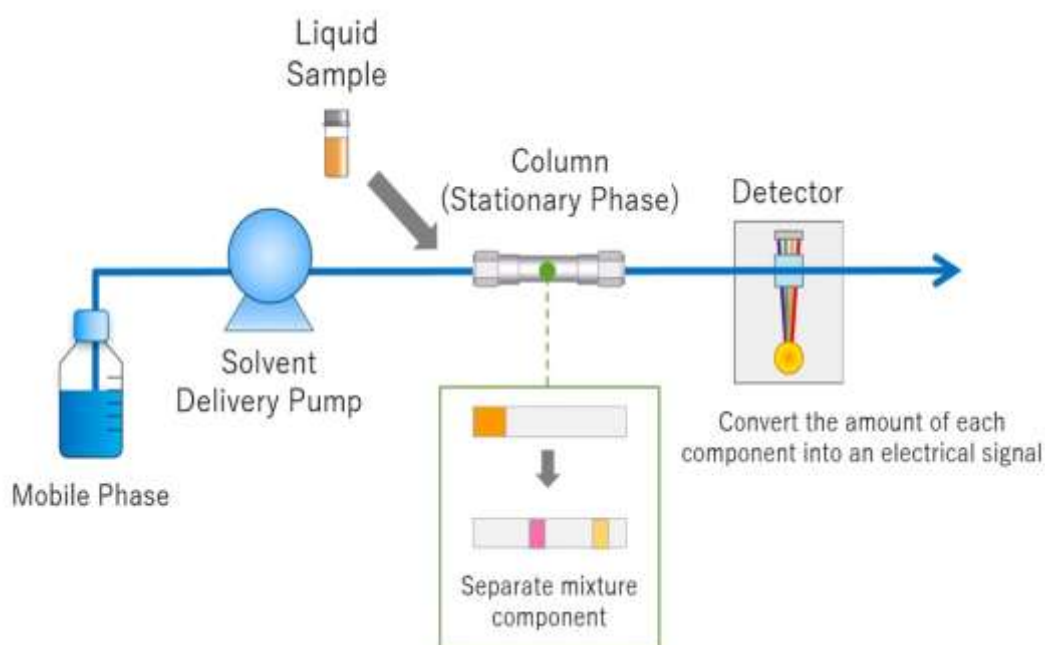


Fig-2 High Performance Liquid Chromatography (HPLC)

In preliminary HPLC (pressure > 20 bar), larger pristine sword columns and stuffing a ccoutrements (flyphspeck size 10- 30 µm are demanded.

The exemplifications of normal phases silica columns are Kromasil 10 µm, Kromasil 16 µm, Chiralcel AS 20 µm whereas for rearphase are Chromasil C18, Chromasil C8, YMC C18. The end is to insulate ordisinfestcomposites, but in logical work the thing is to get in sequence about the sample. This is veritably significant in pharmaceutical assiduity of moment because new p roducts (Natural, Synthetic) have to be introduced to the request as snappily as possible. Having obtainable such an important sanctification fashion makes it possible to spend lower time on the conflation situation².

Liquid Chromatography- Nuclear Magnetic Resonance (LC-NMR)

LC-NMR improves speed and compassion of recognition and found useful in the areas of pharmacokinetics, toxicity studies, drug metabolism and drug detection process. The arrangement of chromatographic separation technique with NMR spectroscopy is one of the most powerful and time saving method for the partition and structural illumination of mysterious compound and mixtures, particularly for the structure clarification of light and oxygen insightful substances. The online LC-NMR technique allows the continuous registration of time changes as they appear in the chromatographic run automated data acquisition and processing in LC- NMR improves speed and sensitivity of detection. The recent prologue of pulsed field gradient technique in high resolution NMR as well as three-dimensional method improves function in structure clarification and molecular weight information. These new hyphenated techniques are useful in the areas of pharmacokinetics, toxicity studies, drug metabolism and drug detection process⁸.

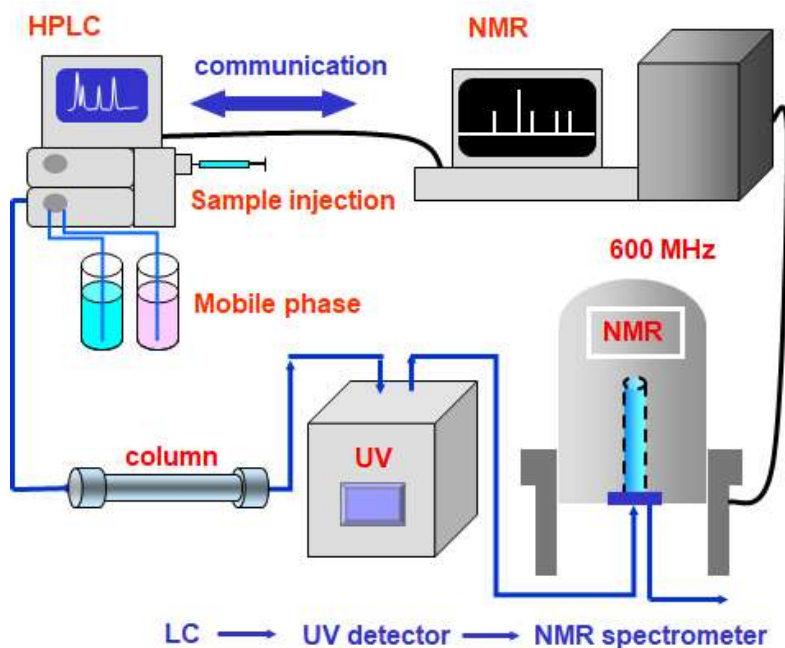


Fig-4 Liquid Chromatography- Nuclear Magnetic Resonance (LC-NMR)

Liquid Chromatography- Mass Spectroscopy (LC-MS)

LC-MS has come system of alternative in numerous stages of medicine expansion. Recent advances includes electro spray, thermo spray, and ion spray ionization ways which offer exclusive advantages o

f high discovery perceptivity and particularity, liquid inferior ion mass spectroscopy, latterly ray mass spectroscopy with 600 MHz offers accurate determination of molecular weight proteins, peptides. Isotopes pattern can be detected by this method⁹.

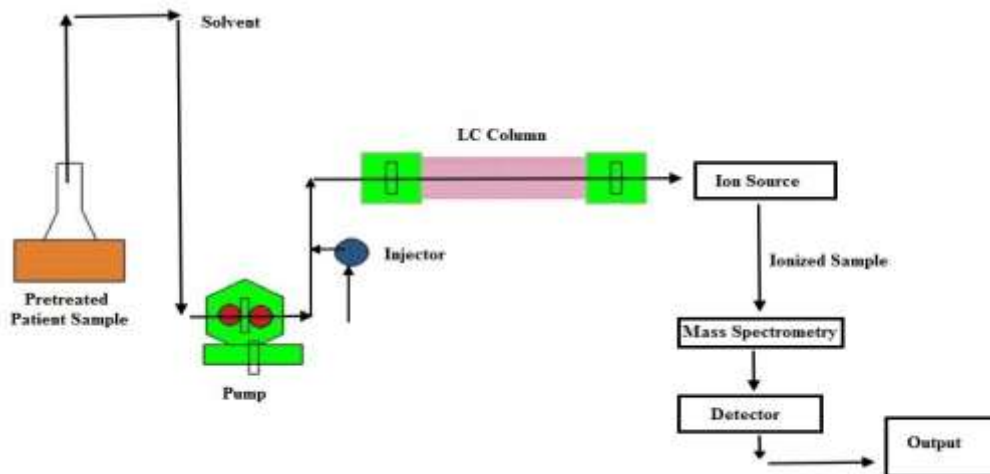


Fig -4 Liquid Chromatography- Mass Spectroscopy (LC-MS)

Nowadays, the most widely utilized bioanalytical method for quantitation is LC-MS/MS, or liquid chromatography (LC) tandem triple-quadrupole mass spectrometry

(MS/MS). The LC/MS/MS procedure is similar to HPLC/UV with regards to sample preparation and chromatographic setup. Compared to UV detection, however, MS/MS offers superior sensitivity and

selectivity, allowing for higher throughput analyses with more sensitive lower quantitation limits. LC/MS/MS is a beneficial, robust, and sensitive procedure used for a wide variety of small molecules. Furthermore, this technology is amenable to automation and unattended analysis.

Gas Chromatography (GC-MS)

It is well identified that numerous pharmacologically active factors in herbal drugs are unpredictable chemical composites. Therefore, the investigation of unpredictable composites by gas chromatography is veritably significant in the investigation of herbal drugs. The GC investigation of the unpredictable canvases has a number of compensations. Originally, the GC of the unpredictable oil painting gives a sensible "point" which can be used to recognize the factory. The composition and virtual attention of the organic composites in the unpredictable oil painting are characteristic of the scrupulous factory and the presence of contaminations in the unpredictable oil painting can be readily detected. Secondly, the birth of the unpredictable oil painting is fairly straightforward and can be formalized and the factors can be readily linked using GC – MS analysis⁹... GC outfit can be straight connived with rapid-fire checkup mass spectrometer of colorful types. GC and GC- MS are generally accepted styles for the analysis of unpredictable ingredients of herbal drugs, due to their perceptivity, constancy and high effectiveness. Particularly, the hyphenation with MS

provides dependable information for the qualitative analysis of the complex ingredients. The inflow rate from the capillary column is usually low sufficient that the article affair can be fed straight into the ionization chamber of MS. The simplest mass sensor in GC is the Ion Trap Sensor (ITD). In this apparatus, ions are twisted from the eluted sample by electron impact or chemical ionization and stored in a radio frequency field; the trapped ions are also ejected from the storehouse area to an electron multiplier sensor. The ejection is illegal so that surveying on the base of mass-to-charge rate is possible. The ions trap sensor is remarkably squashed and less precious than quadruple instruments. GC-MS instruments have been used for recognition of hundreds of factors that are nearby in natural and natural system¹⁰⁻¹¹.

Supercritical Fluid Chromatography (SFC)

Supercritical fluid chromatography is a mongrel of gas and liquid chromatography that combines some of the stylish features of each. SFC permits the separation and determination of a group of composites that aren't accessibly handled by either gas or liquid chromatography. SFC has been applied to a wide variety of accoutrements including natural products, medicines, food and fungicide. These composites are moreover nonvolatile or thermally labile so that GC procedures are irrelevant or contain no functional group that makes possible discovery by the spectroscopic or electrochemical fashion employed in LC¹².

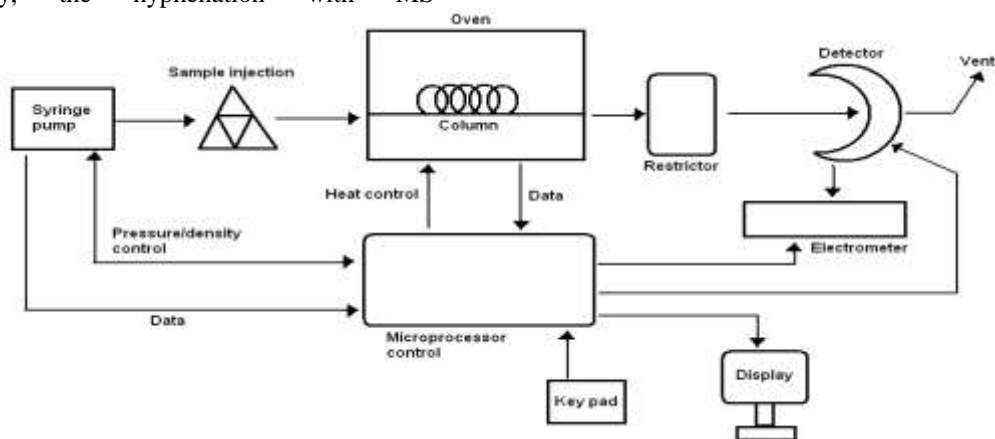


Fig-3 Supercritical Fluid Chromatography (SFC)

II. CONCLUSION

Chromatographic technique is one of the best options for the identification and quantification either in evaluation with chromatography or itself. It is a choice a method for herbal evaluation.

REFERENCE

- [1]. Sim CO, Hamdan MR, Ismail Z and Ahmad MN, Assessment of Herbal Medicines by Chemometrics – Assisted Interpretation of FTIR Spectra, *Journal Of Analytica Chimica Acta*, 2004, 1-14
- [2]. Bhutani KK, Finger-Printing of Ayurvedic Drugs, *The Eastern Pharmacist*, 2000; 507: 21-26.
- [3]. R. Upton, International Symposium on Quality of Traditional Chinese Medicine with Chromatographic Fingerprint, Guangzhou, 2001, i 2-1.
- [4]. Yi-Zeng Lianga, PeishanXieb, Kelvin Chanc, Review: Quality control of herbal medicines; *Journal of Chromatography B*, 812 (2004) 53–70.
- [5]. H. Wagner, S. Bladt, V. Rickl, *Plant Drug Analysis: A Thin Layer Chromatography Atlas*, second ed., Springer-Verlag, 1996.
- [6]. A. Baerheim Svendsen, J. Planar Chromatogr. *Modern TLC 2* (1989) 8.
- [7]. Nikam P. H., Kareparamban J., Jadhav A. and Kadam V., Future Trends in Standardization of Herbal Drugs, *Journal of Applied Pharmaceutical Science* 02 (06); 2012: 38-44.
- [8]. Patil PS, Rajani S. An Advancement of Analytical Techniques in Herbal Research *J. Adv. Sci. Res.* 2010, 1(1); 08-14.
- [9]. Liang YZ, Xie P, Chan K, J., Quality control of herbal medicines, *Chromatogr B*, 2004; 812: 53–70.
- [10]. Guo F.Q., Huang L.F., Zhou S.Y., Zhang T.M., Liang Y.Z., Comparison of the volatile compounds of *Atractylodes* medicinal plants by headspace solid-phase microextraction-gas chromatography–mass spectrometry. *Anal. Chim. Acta* 570: (2006) 73-78
- [11]. Teo C.C., Tan S.N., Yong J.W. H., Hewb C. S., and Ong E. S. Evaluation of the extraction efficiency of thermally labile bioactive compounds in *Gastrodia elata* Blume by pressurized hot water extraction and microwaveassisted extraction. *J. Chromatogr. A* 1182: 2008 34–40.
- [12]. Matthew C, Henry R. Supercritical fluid chromatography, Pressurized liquid extraction, and supercritical fluid extraction. *Anal Chem* 2006; 78: 3909.