ISOLATION AND HIGH-PERFORMANCE THIN-LAYER CHROMATOGRAPHY ANALYSIS OF CONDITIONAL AMINO ACIDS FROM THE FRESH LEAVES OF *ALTERNANTHERA SESSILIS*

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**ABSTRACT**

**Objective:** The objective of the study was to isolate the conditional amino acids present in the fresh leaves extracts of *Alternanthera sessilis* and compare the isolated amino acids with amino acid standards.

**Method:** The fresh plant material was used for extraction. For the extracted sample solution, paper chromatography and high-performance thin-layer chromatography (HPTLC) fingerprinting analysis were then carried out.

**Results:** Since *A. sessilis* is rich in protein, various amino acids have been isolated in aggregates and paper chromatography has been performed. Amino acids were separated and identified using trial and error method. Cysteine, glycine, aspartic acid, and proline are the standard amino acids used for comparing with the isolated amino acids by HPTLC fingerprint analysis, which gives an idea for the authentication of the plant, its constituents and also provides a parameter for quality control.

**Conclusion:** The results scientifically validate the use of *A. sessilis* in the traditional medicine and also as a food.

**Keywords:** Fresh leaves extract, *Alternanthera sessilis*, Amino acids, High-performance thin-layer chromatography, Traditional medicine.
Herbs are usually rich in amino acid, one such plant is *A. sessilis*, and its phytochemical analysis reveals the presence of amino acid [5] and also possesses wound healing activity [6]. *A. sessilis* commonly known as sessile joyweed is a perennial herb, often found in and near ponds, canals, and reservoirs reported 5% of protein. In traditional medicine, the whole plant is used internally and externally and used to treat diarrhea, skin diseases, night blindness, indigestion, and fever. *A. sessilis* is being widely used for cooking by people of Tamil Nadu and other Southern states of India.

**METHODS**

**Collection of plant material**

*A. sessilis* leaves were collected in and around in Coimbatore District, Tamil Nadu, India, and authenticated by Dr. C Murugan, Scientist “D” and Head of office, TN.AU, Coimbatore, Tamil Nadu, India. Authentication number BSI/SRC/5/23/2017/Tech-3527.

**Isolation of free amino acid**

Fresh leaves (150 mg) of *A. sessilis* were collected and kept in a frozen condition along with the liquid nitrogen to prevent degradation. Frozen leaves were grinded in a dry mortar along with liquid nitrogen and 600 μl of a mixture of solutions containing water, chloroform, and methanol in a ratio (3:5:12 v/v). Addition of methanol makes the mixture homogenized. This homogenate is completely transferred to 2 ml centrifuge tube and remaining in the mortar was also rinsed with mixture of solution and transferred to the centrifuge. Immediately the mixture was centrifuged at full speed for 2 min and supernatant solution was collected separately.

Re-extract the residue with another 600 μl of mixture of solutions containing water, chloroform, and methanol for about 2 min and only the supernatant solution were transferred to fresh tube. Centrifugation continued till clear solution is obtained. Samples were stored at −80°C and the extracted amino acids were quantified using ninhydrin assay. Usually, amino acid will be degraded within 1 week, so it is being quantified immediately [7].

This method is more reliable and less costly. Hence, free amino acid can be isolated from the fresh leaves of *A. sessilis* and can be used for further work.

**Paper chromatography**

Paper chromatography is an analytical method used to separate colored chemicals or substances. In paper chromatography, the sample mixture is applied to a piece of Whatman filter paper using the capillary tube and the edge of the paper is immersed in a solvent (mobile phase) and the solvent moves up the paper by capillary action. The filter paper is then dried and sprayed with ninhydrin reagent. Once again stationary phase dried in a hot air oven for 110°C.

Amino acids isolated from fresh leaves are separated using chromatogram and compared with various standard amino acids using trial and error method [8].

**RESULTS AND DISCUSSION**

**Isolation of free amino acid**

From fresh leaves juice of *A. sessilis*, as per the procedure discussed earlier various free amino acids are isolated in the form of aggregates. Hence, to separate the amino acids aggregates and also to assume the amino acid, paper chromatography method is applied.

**Paper chromatography**

Since paper chromatography is one of the reliable methods for determination of amino acid, isolated amino acids are identified using trial and error method. n-Butanol:glacial acetic acid:water in a ratio 4:5:1 is used as a mobile phase since polar side chain amino acids (aspartic acid, cysteine, lysine, etc.) and shorter non-polar amino acids (proline, alanine, and glycine) migrates faster in this solvent. The sample is separately developed and viewed at 362 nm reported in Fig. 1 and also viewed after spraying detecting agent (ninhydrin reagent) is reported in Fig. 2.

Then using trial and error method, the sample containing amino acid is being compared with various standard amino acids and four amino acids have been assumed based on the coinciding R values.
The amino acids assumed by paper chromatography are cysteine, glycine, aspartic acid, and proline.

**HPTLC analysis of amino acid**

Seven different types of amino acids were observed with peak value from the isolated sample of *A. sessilis*. The suitable solvent system selected for evaluation was n-Butanol:glacial acetic acid:water (4:1:5).

The chromatogram of the sample was developed along with the four standard amino acids. HPTLC fingerprint of a plant species provides basic information regarding isolation, purification, and characterization. This present study provides sufficient information about amino acids present in fresh leaves of *A. sessilis* and also identification, standardization and quality control of the medicinal plant. HPTLC chromatogram of the sample was reported in Table 1, indicates the presence of seven compounds.

- **Fig. 1**: Paper chromatography of isolated amino acid detected at UV 362nm
- **Fig. 2**: Paper chromatography of isolated amino acid detected using ninhydrin
- **Fig. 3**: 3-Dimensional view of chromatogram of sample with four standard amino acids
- **Fig. 4**: High-performance thin-layer chromatography chromatogram of the amino acid at 366 nm
- **Fig. 5**: High-performance thin-layer chromatography chromatogram of the amino acid at 254 nm

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- **Fig. 1**: Paper chromatography of isolated amino acid detected at UV 362nm
- **Fig. 2**: Paper chromatography of isolated amino acid detected using ninhydrin
- **Fig. 3**: 3-Dimensional view of chromatogram of sample with four standard amino acids
- **Fig. 4**: High-performance thin-layer chromatography chromatogram of the amino acid at 366 nm
- **Fig. 5**: High-performance thin-layer chromatography chromatogram of the amino acid at 254 nm

The results from HPTLC scanned at 366 nm and reported in Fig. 4, for the amino acid isolated from fresh leaves of *A. sessilis*, reveals the presence of 5 amino acids with the *R* values: 0.35, 0.43, 0.55, 0.72, and 0.77. The *R* values for standard amino acids cysteine, glycine, aspartic acid, and proline are 0.21, 0.39, 0.37, and 0.40, respectively. Comparing with the four standard amino acids, the *R* value of standard (I) 0.37 coincide with *R* value of sample 0.35 and the *R* value of standard (II) 0.40 coincide with *R* value of sample 0.43.

The results from HPTLC scanned at 254 nm and reported in Fig. 5, for an isolated sample containing amino acid reveals the presence of 6 amino acids with the *R* values: 0.03, 0.20, 0.35, 0.46, 0.56, and 0.74. The *R* values for standard amino acids cysteine, glycine, aspartic acid, and proline are 0.20, 0.40, 0.37, and 0.41, respectively. When comparing with the four standard amino acids, the *R* value of standard (I) 0.20 coincide with *R* value of sample 0.20, the *R* value of standard (II) 0.40 coincide with *R* value of sample 0.35, and the *R* value of standard (III) 0.41 coincide with *R* value of sample 0.46.
HPTLC chromatogram was developed; ninhydrin reagent was sprayed and observed. HPTLC fingerprint sample has been compared with standard amino acids - aspartic acid, glycine, cysteine, and proline and reported in Figs. 6-9.

**Standard I** - Cysteine  
**Standard II** - Glycine  
**Standard III** - Aspartic acid  
**Standard IV** - Proline.

**DISCUSSION**

Mixture of amino acids has been successfully isolated from the fresh leaves of *A. sessilis* and assumed using paper chromatogram (trial and error method). Based on the paper chromatogram, standard amino acids are decided. From HPTLC, we can conclude the presence of amino acids such as cysteine, aspartic acid, and proline. Hence, plant or plant extract can be given in deficiency of these conditional amino acids.
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The result obtained by qualitative evaluation of HPTLC will be helpful in the further quantitative estimation of each and every amino acid and also for standardization of herbal formulations containing this plant.

CONCLUSION

It can be concluded that based on the results obtained from the HPTLC fingerprint analysis will be helpful in identification and standardization of *A. sessilis* and can be utilized as a reference for the identification and quality control of the drug. The present study can be considered as reference and can be used for validation of the plant usage.

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AUTHORS’ CONTRIBUTION

All authors have contributed equally to carry out the work.

CONFLICTS OF INTEREST

The authors declared that they have no conflicts of interest.

REFERENCES


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**Table 1: HPTLC profile for sample**

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HPTLC: High-performance thin-layer chromatography.